

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: April 19, 2005, 15:35:11 ; Search time 43.7895 Seconds

(without alignments)
1413.163 Million cell updates/secTitle: US-09-904-954-4_COPY_1_160
Perfect score: 879
Sequence: 1 MRLPLRLRTVLMMAFLGSP...SGQCLRLQVSVCKERSSES 160Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 1

Minimum DB seq length: 0
Maximum DB seq length: 2000000000Post-processing: Minimum Match 89%
Maximum Match 100%
Listing first 1500 summariesDatabase : A_Geneseq_16Dec04:*
1: geneseqp1980s:*
2: geneseqp1990s:*
3: geneseqp2000s:*
4: geneseqp2001s:*
5: geneseqp2002s:*
6: geneseqp2003as:*
7: geneseqp2003bs:*
8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	879	100.0	201	2 AAR71482	Aar71482 Human hek

ALIGNMENTS

RESULT 1
ID AAR71482 standard; protein; 201 AA.
XX
AC AAR71482;
XX
DT 25-MAR-2003 (revised)
DT 03-OCT-1995 (first entry)
XX
DE Human hek-L protein.
XX
KW Ligand; cell surface; tyrosine kinase receptor; tumorigenesis; immunogen.
XX
OS Homo sapiens.
XX
FH Key
FH Peptide
FT 1..22
FT /note= "signal peptide"
FT Protein 23..201

XX WO9506065-A1.
XX 02-MAR-1995.
XX 17-AUG-1994; 94WO-US009282.
XX 20-AUG-1993; 93US-00109745.
XX 30-AUG-1993; 93US-00114426.
XX 03-DEC-1993; 93US-00161132.
XX 09-MAY-1994; 94US-00240124.
XX (IMMV) IMMUNEX CORP.
XX Beckmann MP, Cerretti DP;
XX WPI; 1995-106811/14.
XX N-PSDB; AAQ85888.
XX New isolated DNA encoding hek-L protein or its fusion products - useful
XX as assay reagent or for carrying therapeutic and diagnostic compounds to
XX leukaemia cells.
XX Claim 21; Page 38; 45pp; English.
XX The sequence is that of a novel protein designated hek-L, a protein that
XX can bind hek (a cell surface receptor tyrosine kinase). Hek-L is the
XX first known ligand for hek and can be used to study cellular processes
XX regulated by hek (which may be involved in tumorigenesis). It is also an
XX immunogen for antibody production, as a reagent for detecting hek or hek-
XX L in vitro assays, to determine binding of hek proteins, to purify hek
XX proteins, and to carry diagnostic or cytotoxic agents to particular
XX leukaemia cells that express the hek antigen. Hek-L also binds the alk
XX tyrosine kinase receptors. See also AAR71481. (Updated on 25-MAR-2003 to
XX correct PN field.)
XX Sequence 201 AA:

Query Match 100.0%; Score 879; DB 2; Length 201;
Best Local Similarity 100.0%; Pred. No. 3.4e-93;
Matches 160; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MRLPLRLRTVLMMAFLGSP...SGQCLRLQVSVCKERSSES 160
DB 1 MRLPLRLRTVLMMAFLGSP...SGQCLRLQVSVCKERSSES 160
QY 61 YEGPGPEGEPTFALVMDWPGYESSCOAEGPRAYKRWVCSLPFGHVQFSEKIQRTFTFSL 120
DB 61 YEGPGPEGEPTFALVMDWPGYESSCOAEGPRAYKRWVCSLPFGHVQFSEKIQRTFTFSL 120
QY 121 GFEFLPGETYYIISVPTPESSGQCLRLQVSVCKERSSES 160
DB 121 GFEFLPGETYYIISVPTPESSGQCLRLQVSVCKERSSES 160

Search completed: April 19, 2005, 23:13:33
Job time : 43.7895 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 19, 2005, 09:30:04 ; Search time 3056 Seconds
(without alignments)
8183.329 Million cell updates/sec

Title: US-09-904-954-1_COPY_140_796
Perfect score: 657
Sequence: 1 CTGCTGGCCCAAGGAGGCGCCG.....TCATGACGTTCTTGCCCTCC 657

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 5

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 89%
Maximum Match 100%
Listing first 1500 summaries

Database :

EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_hnc:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_g881:*
9: gb_g882:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	657	100.0	1741	3	CR607482 full-leng
2	645	98.2	871	5	BX419695 BX419695
3	622.6	94.8	1100	1	AL533153 AL533153
4	598.4	91.1	726	1	AL527972 AL527972
5	587.2	89.4	1629	3	CR597504 CR597504 full-leng

ALIGNMENTS

RESULT 1
CR607482
LOCUS CR607482 1741 bp mRNA linear HTC 21-JUL-2004
DEFINITION full-length cDNA clone CSODN03YL19 of Adult brain of Homo sapiens (human).
ACCESSION CR607482
VERSION CR607482.1 GI:50488289
KEYWORDS HTC; CNSLT cDNA
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 1741)
AUTHORS Li, W. B., Gruber, C., Jesse, J. and Polayes, D.

TITLE Full-length cDNA libraries and normalization
JOURNAL Unpublished
REMARK Contact: Feng Liang Email: fliang@lifetech.com URL: http://fulllength.invitrogen.com/Invitrogen Corporation 1600 Faraday Avenue
2 (bases 1 to 1741)
Genoscope.

REFERENCE
AUTHORS Direct Submission
JOURNAL Submitted (20-JUL-2004) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail: seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-stranded cDNA was digested with Not I and cloned into the Not I and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies, a division of Invitrogen.
Location/Qualifiers
1. 1741
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CSODN03YL19"
/issue_type="Adult brain"
/plasmid="pCMVSPORT_6"

COMMENT

FEATURES
source

ORIGIN

Query Match 100.0%; Score 657; DB 3; Length 1741;
Best Local Similarity 100.0%; Pred. No. 6.2e-143;
Matches 657; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	CTGCTGGCCCAAGGAGGCGCCGCTGGGAAACCGGATGGGCTGTAAGACG 60
DB	122	CTGCTGGCCCAAGGAGGCGCCGCTGGGAAACCGGATGGGCTGTAAGACG 181
QY	61	TCCACACGACCTGCGGCGAGAGGCTACACCGTGAAGCTGAACGATATCTG 120
DB	182	TCCACACGACCTGCGGCGAGAGGCTACACCGTGAAGCTGAACGATATCTG 241
QY	121	GATTTTATCTGCGCGCACTACAAAGCTCGGGGTGGGCCCGGCGGAGCGGCGCC 180
DB	242	GATTTTATCTGCGCGCACTACAAAGCTCGGGGTGGGCCCGGCGGAGCGGCGCC 301
QY	181	GGAGCGGGGCGAGAGCTAGTGTGTATATGTTGAGCGGCAACCGCTACCGACCTGC 240
DB	302	GGAGCGGGGCGAGAGCTAGTGTGTATATGTTGAGCGGCAACCGCTACCGACCTGC 361
QY	241	AACGCCACGAGGCTTCAAGCGCTGGAGTGAACCGCGGACGCGCCGACAGCGCC 300
DB	362	AACGCCACGAGGCTTCAAGCGCTGGAGTGAACCGCGGACGCGCCGACAGCGCC 421
QY	301	ATCAAGTTCTCGAGAAAGTTCCAGCGCTACAGCGCTTCTCTGGGCTACGAGTTCCAC 360
DB	422	ATCAAGTTCTCGAGAAAGTTCCAGCGCTACAGCGCTTCTCTGGGCTACGAGTTCCAC 481
QY	361	GCGGCGGAGAGTCTATCTATCTCAAGCCCACTCAAACTGCACTGGAATGCTCTG 420
DB	482	GCGGCGGAGAGTCTATCTATCTCAAGCCCACTCAAACTGCACTGGAATGCTCTG 541
QY	421	AGGATGAAGTGTTCGCTGCTGCGCTCCACATCGCACTCCGGGAGAGCGGATCCCC 480
DB	542	AGGATGAAGTGTTCGCTGCTGCGCTCCACATCGCACTCCGGGAGAGCGGATCCCC 601
QY	481	ACTCTCCCGCAGTTACCATGAGGCGCCCAATGTGAAGATCAAGTGTGGAAGCTTTGAG 540
DB	602	ACTCTCCCGCAGTTACCATGAGGCGCCCAATGTGAAGATCAAGTGTGGAAGCTTTGAG 661
QY	541	GGAGAGAACCTTAAGTGGCCCAAGCTTGAAGAAGATCAAGGAGCGGAGCGGCAACCGG 600
DB	662	GGAGAGAACCTTAAGTGGCCCAAGCTTGAAGAAGATCAAGGAGCGGAGCGGCAACCGG 721
QY	601	GAACACCTGCGCTGGCGGCTGAGCATGCGCTTCTCTCAATGACGTTCTTGAGCTCC 657
DB	722	GAACACCTGCGCTGGCGGCTGAGCATGCGCTTCTCTCAATGACGTTCTTGAGCTCC 778

RESULT 2
LOCUS BX419695
DEFINITION BX419695 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone
ACCESSION BX419695
VERSION BX419695.2
KEYWORDS GI:46934231
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1. W.B., Gruber, C., Jessee, J. and Polyes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
JOURNAL On May 13, 2003 this sequence version replaced gi:30654816.
COMMENT Contact: Genoscope
Genoscope - Centre National de Sequencage
2 rue Gaston Cremieux, CP 5706 - 91057 EVRY cedex - FRANCE
Email: seqref@genoscope.cns.fr; Web: www.genoscope.cns.fr
1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime
into the Not I and EcoRV sites of the pCMVSPORT 6 vector. Library
was not normalized. Library was constructed by Life Technologies, a
division of Invitrogen.
This sequence belongs to sequence cluster 497.f
For more information about this cluster, see
http://www.genoscope.cns.fr/cdna?c=CS0DF019D1A20P1&c=497.f

FEATURES
Source
1. 871
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CS0DF019YB24"
/tissue_type="FETAL BRAIN"
/dev_stage="fetal"
/clone_lib="Homo sapiens FETAL BRAIN"
/note="Organ: brain; Vector: pCMVSPORT_6; 1st strand cDNA
enriched, double-strand cDNA was digested with Not I and
cloned into the Not I and EcoRV sites of the pCMVSPORT 6
vector. Library was not normalized."

Query Match
Best Local Similarity 98.2%; Score 645; DB 5; Length 871;
Matches 656; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY 1 CTGCTGCCCAAGGCGCGAGGCGCTGGGAAACCGGCGATGCGTGTACTGGAACAGC 60
DB 114 CTGCTGCCCAAGGCGCGAGGCGCTGGGAAACCGGCGATGCGTGTACTGGAACAGC 60
QY 61 TCCAACCAAGCACTGCGCGAGGCGCTGACCGTGAAGTGAACGACTATCTG 120
DB 173 TCCAACCAAGCACTGCGCGAGGCGCTGACCGTGAAGTGAACGACTATCTG 120
QY 121 GATATTTATGCGCGCGAGGCGCTGACCGTGAAGTGAACGACTATCTG 232
DB 233 GATATTTATGCGCGCGAGGCGCTGACCGTGAAGTGAACGACTATCTG 232
QY 181 GGAGGCGGCGAGGCGAGTACGTGCTGAACCTGTGAGCGCAAGCGCTACCTG 240
DB 293 GGAGGCGGCGAGGCGAGTACGTGCTGAACCTGTGAGCGCAAGCGCTACCTG 240
QY 241 AACGCGAGGCGAGGCGTGTGAGGAGTGAACCGGCGCGACCGCTGCAAGCTG 352
DB 353 AACGCGAGGCGAGGCGTGTGAGGAGTGAACCGGCGCGACCGCTGCAAGCTG 352
QY 301 ATCAAGTCTGCGAGGAGTTCAGCGCTGACCGCTTCTCTGCTGAGTCTGAC 360
DB 412 ATCAAGTCTGCGAGGAGTTCAGCGCTGACCGCTTCTCTGCTGAGTCTGAC 360

RESULT 3
LOCUS AL533153
DEFINITION AL533153 Homo sapiens ADULT BRAIN Homo sapiens cDNA clone
ACCESSION CS0DN003YD19 5-PRIME, mRNA sequence.
VERSION AL533153.3
KEYWORDS GI:45708058
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1. W.B., Gruber, C., Jessee, J. and Polyes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
JOURNAL On Feb 13, 2001 this sequence version replaced gi:31070985.
COMMENT Contact: Genoscope
Genoscope - Centre National de Sequencage
2 rue Gaston Cremieux, CP 5706 - 91057 EVRY cedex - FRANCE
Email: seqref@genoscope.cns.fr; Web: www.genoscope.cns.fr
1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime
into the Not I and EcoRV sites of the pCMVSPORT 6 vector. Library
was not normalized. Library was constructed by Life Technologies, a
division of Invitrogen.
This sequence belongs to sequence cluster 497.f
For more information about this cluster, see
http://www.genoscope.cns.fr/cdna?c=CS0DN003YD19P1&c=497.f

FEATURES
Source
1. 1100
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CS0DN003YD19"
/tissue_type="ADULT BRAIN"
/dev_stage="adult"
/clone_lib="Homo sapiens ADULT BRAIN"
/note="Organ: brain; Vector: pCMVSPORT_6; 1st strand cDNA
enriched, double-strand cDNA was digested with Not I and
cloned into the Not I and EcoRV sites of the pCMVSPORT 6
vector. Library was not normalized."

Query Match
Best Local Similarity 99.8%; Score 622.6; DB 1; Length 1100;
Matches 655; Conservative 1; Mismatches 0; Indels 3; Gaps 3;
QY 1 CTGCTGCCCAAGGCGCGAGGCGCTGGGAAACCGGCGATGCGTGTACTGGAACAGC 60

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Db      122 CTGCTGAGCCCAAGAGCCCGGA-GGGCCCTGGGAAACCGGATGGGCTGATGGAACAGC 180
Qy      61  TCACACCAAGACCTTCGGCGGAGAGGCTACACCGCTGACAGTGAACGATATCTG 120
Db      181  TCCAAACAGACCTCGCGCGAGAGGCTACACCGCTGACAGTGAACGATATCTG 240
Qy      121  GATATTACTGCGCGCACTACAAAGCTCGGGGGTGGGGCCCGGGGGGGGCGGGGGCC 180
Db      241  GATATTACTGCGCGCACTACAAAGCTCGGGGGTGGGGCCCGGGGGGGGCGGGGGCC 300
Qy      181  GAGAGCGGGGAGAGAGAGTACGTCTGATACGTGAGCCGCAACGCTACCGGACCTGC 240
Db      301  GAGAGCGGGGAGAGAGAGTACGTCTGATACGTGAGCCGCAACGCTACCGGACCTGC 360
Qy      241  AAGCCACGACGAGGCTTCAAGCGCTGGGAGTGAACCGGCGGACGCGCCGACAGCCCC 420
Db      361  AAGCCACGACGAGGCTTCAAGCGCTGGGAGTGAACCGGCGGACGCGCCGACAGCCCC 420
Qy      301  ATCAAGTTCTCGAGAGAGTTCAGCGCTACGCGGCTTCTCT-GGGCTACGAGTTCCA 359
Db      421  ATCAAGTTCTCGAGAGAGTTCAGCGCTACGCGGCTTCTCTCTGCGGCTACGAGTTCCA 480
Qy      360  CGCGGCGCAGAGTACTACTACATCTCCAGCCGCTACCAACCTGCACTGGAAGTGTCT 419
Db      481  CGCGGCGCAGAGTACTACTACATCTCCAGCCGCTACCAACCTGCACTGGAAGTGTCT 540
Qy      420  GAGAGTGAAGGTTCTGCTGCTGCGGCTTCACATGCACTCC-GGGAGAGAGCGGTC 478
Db      541  GAGAGTGAAGGTTCTGCTGCTGCGGCTTCACATGCACTCCGCGGAGAGAGCGGTC 600
Qy      479  CCACTCTCCCGCAGTTACCATATGGGCCCCCAATGGAAGATCAAGTGTCTGGAACCTTTG 538
Db      601  CCACTCTCCCGCAGTTACCATATGGGCCCCCAATGGAAGATCAAGTGTCTGGAACCTTTG 660
Qy      539  AGGAGAGAACCTTCAGAGTGGCCCAAGCTTGAGAGAGATCAGGCGGAGCCGCAAC 598
Db      661  AGGAGAGAACCTTCAGAGTGGCCCAAGCTTGAGAGAGATCAGGCGGAGCCGCAAC 720
Qy      599  GGGAAACACTGCGCCCTGCGCGTGGGCAATCGCTTCTCTCATGACGTTTGGCCTCC 657
Db      721  GGGAAACACTGCGCCCTGCGCGTGGGCAATCGCTTCTCTCATGACGTTTGGCCTCC 779

```

RESULT 4
 AL527972
 LOCUS
 DEFINITION
 AL527972 Homo sapiens NEUROBLASTOMA COT 25-NORMALIZED Homo sapiens
 cDNA clone CS0DC027YP19 5-PRIME, mRNA sequence.
 AL527972
 VERSION
 AL527972.3 GI:45703058
 KEYWORDS
 EST.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 726)
 Full-length cDNA libraries and normalization
 Unpublished (2001)
 On Feb 13, 2001 this sequence version replaced gi:31065823.
 COMMENT
 Contact: Genoscope
 Genoscope - Centre National de Sequencage
 2 rue Gaston Cremieux, CP 5706 - 91057 Evry cedex - FRANCE
 Email: seqref@genoscope.cns.fr, Web: www.genoscope.cns.fr
 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime
 into the NotI and EcoRV sites of the pCMVSPORT 6 vector. Library
 was normalized. Library was constructed by Life Technologies, a
 division of Invitrogen. This sequence belongs to sequence cluster
 497.f
 For more information about this cluster, see
 http://www.genoscope.cns.fr/cdnats=CS0DC027CH10P1&c=497.f.

FEATURES

source

Location/Qualifiers
 1..726
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="CS0DC027YP19"
 /issue_type="NEUROBLASTOMA COT 25-NORMALIZED"
 /clone_lib="Homo sapiens NEUROBLASTOMA COT 25-NORMALIZED"
 /note="1st strand cDNA was primed with a NotI-oligo(dT)
 primer. Five prime end enriched, double-strand cDNA was
 digested with NotI and cloned into the NotI and EcoRV
 sites of the pCMVSPORT 6 vector. Library was normalized."

ORIGIN

Query Match

91.1%; Score 598.4; DB 1; Length 726;

Best Local Similarity

91.0%; Pred. No. 2,86-129;

Matches 595; Conservative 41; Mismatches 17; Indels 1; Gaps 1;

```

Qy      1  CTGCTGAGCCCAAGAGGCGGCTGGGAAACCGGATGGGCTGATGGAACAGC 60
Db      69  CTGCTGAGCCCAAGAGGCGGCGA-GGGCGAAGGAAACGAGATGGGCTGATGGAACAGC 127
Qy      61  TCACACCAAGACCTTCGGCGGAGAGGCTACACCGCTGACAGTGAACGATATCTG 120
Db      128  WCCAAACAGACCTCGCGCGAGAGGCGCMAAMMAAGAGTGAACGTAACGATATCTG 187
Qy      121  GATATTACTGCGCGCACTACAAAGCTCGGGGGTGGGGCCCGGGGGGGGAGCCGCGGCC 180
Db      188  GATATMMAAAGCAGMAMTAAAGCMCGGGGTRGGCCCGGGGGGAGMCGGGGCC 247
Qy      181  GAGAGCGGGGAGAGAGTACGTCTGATACGTGAGCCGCAACGAGCTACCGGACCTGC 240
Db      248  GAGAGCGGGGAGAGAGTACGTCTGATACGTGAGCCGCAACGAGCTACCGGACCTGC 307
Qy      241  AAGCCACGACGAGGCTTCAAGCGCTGGGAGTGAACCGGCGGACGCGCCGCAACGAGCC 300
Db      308  AAGCCACGACGAGGCTTMAAGCGTGGAGATGGAACCGGCGGACGCGCCGCAACGAGCC 367
Qy      301  ATCAAGTTCTCGAGAGAGTTCAGCGCTACAGCGCTTCTCTCTGCGGCTACGAGTTCCAC 360
Db      368  ATMAAGTTCTCGAGAGAGTTCAGCGCGMAAARCGCTTCTCTCTGCGGCGMAAGATGCCAC 427
Qy      361  GCGGCGCAGAGTACTACTACATCTCCAGCCGCTTCACATGCACTCCGGAAGTGTCTG 420
Db      428  GCGGCGCAGAGTACTACTACATCTCCAGCCGCTTCACATGCACTCCGGAAGTGTCTG 487
Qy      421  AGATGAAGGTTCTGCTGCTGCGGCTTCACATGCACTTCGCGGAGAGCCGCTCCC 480
Db      488  AGATGAAGGTTCTGCTGCTGCGGCTTCACATGCACTTCGCGGAGAGCCGCTCCC 547
Qy      481  ACTCTCCCGCAGTTACCATATGGGCCCCCAATGGAAGATCAAGTGTCTGGAAGCTTTGAG 540
Db      548  ACTCTCCCGCAGTTACCATATGGGCCCCCAATGGAAGATCAAGTGTCTGGAAGCTTTGAG 607
Qy      541  GAGAGAGAACCTTCAGAGTGGCCCAAGCTTGAGAGAGATCAGGCGGAGCCGCAACG 600
Db      608  GAGAGAGAACCTTCAGAGTGGCCCAAGCTTGAGAGAGATCAGGCGGAGCCGCAACG 667
Qy      601  GAACACCTGCGCCCTGCGCGTGGGCAATCGCTTCTCTCATGACGTTTGGCCTCC 654
Db      668  GAACACCTGCGCCCTGCGCGTGGGCAATCGCTTCTCTCTCATGACGATATCTTGGC 721

```

RESULT 5
 CRS97504
 LOCUS
 DEFINITION
 CRS97504 1629 bp mRNA linear HTC 21-JUN-2004
 full-length cDNA clone CS0D1026T124 of Placenta Cot 25-normalized
 of Homo sapiens (human).
 ACCESSION
 CRS97504
 VERSION
 CRS97504.1 GI:50478311
 KEYWORDS
 HTC; CNSLT; cDNA.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Homo sapiens

us-09-904-954-1_copy_140_796.rst

AUTHORS Li, W. B., Guber, C., Jesse, J. and Polayes, D.
TITLE Full-length cDNA libraries and normalization
JOURNAL Unpublished
REMARK Contact : Feng

COMMENT

ORIGIN

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/mol_type="rRNA"
/db_xref="taxon:9606"
/clone="CS0D1026Y124"
/tissue_type="Placenta"
/plasmid="pCMVSPORT_6"
Cot 25-normalized

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Query Match	89.4%	Score 587.2	DB 3	Length 1629
Best Local Similarity	99.5%	Pred. No. 1,3e-12;		
Matches 589; Conservative	0	Mismatches		

16 CCAGACCTGGCGGAGAGGGCTAACCCGTGAGGTGAACCTGAACCACTATCTGGATAT 125
Db |||
77 CCACAGCCTGGCGGAGAGGGCTAACCGTGCAGTGAAGTGAACCACTATCTGGATAT 125
Qy |||
126 TTACTGGCCCCACTTACACAGCTCCGGGGTGGGCCCCCGGGCGGGAACCGGGCCCGGAGG 136
Db |||
137 TTACTGCCGCACTACACAGCTGGGGGTGGGCCCCCGGGCGGGAACCGGGCCCGGAGG 185
Qy |||
186 CCGGGCAGAGCAGTACGTGCTGTACATGTGAACCGCAACCGCTACCGCACCTGCAACGC 196
Db |||
197 CCGGGCAGAGCAGTACGTGCTGTACATGTGAACCGCAACCGCTACCGCACCTGCAACGC 245
Qy |||
246 CAGCCAGGGCTTCAAGAGGCTGGGAGTGCACACCGGCGGACCGCCGCAAGGCCCATCA 256
Db |||
257 CAGCCAGGGCTTCAAGAGGCTGGGAGTGCACACCGGCGGACCGCCGCAAGGCCCATCA 305
Qy |||
306 GTTCTCGGAGAGTTCACGCGTTACAGGCGCTTCTCTCTGGCTACGAGTTCACGCGCG 316
Db |||
317 GTTCTCGGAGAGTTCACGCGTTACAGGCGCTTCTCTCTGGCTACGAGTTCACGCGCG 365
Qy |||
366 CCAAGAGTACTTACATCTCCACGCCCACTCACCACTGCACTGGAAGTGCAGAGAT 376
Db |||
377 CCAAGAGTACTTACATCTCCACGCCCACTCACCACTGCACTGGAAGTGCAGAGAT 425
Qy |||
426 GAAAGTGTTCGTCTGCTCGCGCTCCACATGCACTCCGGGGAGAAAGCCGGTCCCACTCT 436
Db |||
437 GAAAGTGTTCGTCTGCTCGCGCTCCACATGCACTCCGGGGAGAAAGCCGGTCCCACTCT 485
Qy |||
486 CCCCAGTTCAACATGGGCCCCCAATGTGAAGATCAAGCTGTGGAGAATCTTTGAGGGAGA 496
Db |||
497 CCCCAGTTCAACATGGGCCCCCAATGTGAAGATCAAGCTGTGGAGAATCTTTGAGGGAGA 545
Qy |||
546 GAAACCTCAGAGTCCCAAGCTTGAAGAGAGATAGAGGGGACCAAGCCCAACCGGAAACA 556
Db |||
557 GAAACCTCAGAGTCCCAAGCTTGAAGAGAGATAGAGGGGACCAAGCCCAACCGGAAACA 605
Qy |||
606 CCGGCCCCCTGGCCGTGGGCATCGCTTCTCTCATGACGTTCTTGGCGTCC 657
Db |||
617 CCGGCCCCCTGGCCGTGGGCATCGCTTCTCTCATGACGTTCTTGGCGTCC 668

Search completed: April 19, 2005, 21:57:08
Job time : 3057.14 secs

Wed Apr 20 07:32:03 2005

QY 661 CAC 663
Db 671 CAC 673

Search completed: April 19, 2005, 21:57:07
Job time : 3085.05 secs

us-09-904-954-1_copy_83_745.rst

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: April 19, 2005, 15:35:11 ; Search time 59.9368 Seconds
(without alignments)
1413.163 Million cell updates/sec

Title: US-09-904-954-2_COPY_1_219

Perfect score: 1207
Sequence: 1 MAAAPLLLLLVLPVPLPLP.....GENPQVKLEKISIGTSFKR 219

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 6

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 89%
Maximum Match 100%
Listing first 1500 summaries

Database : A_Geneseq_16Dec04:*
1: geneseqp1980s:*
2: geneseqp1990s:*
3: geneseqp2000s:*
4: geneseqp2001s:*
5: geneseqp2002s:*
6: geneseqp2003as:*
7: geneseqp2003bs:*
8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1207	100.0	238	2 AAR71481	AAR71481 Human hek
2	1207	100.0	238	5 AAG79439	AAG79439 Breast/co
3	1207	100.0	238	6 ABP97190	ABP97190 Tumour/as
4	1207	100.0	238	7 ADN38812	ADN38812 Cancer/an
5	1207	100.0	238	8 ADC21666	ADC21666 Human sof
6	1172	97.1	234	2 AAR82605	AAR82605 Eph trans

ALIGNMENTS

RESULT 1
ID AAR71481 standard; protein; 238 AA.
XX
AC AAR71481;
XX
DT 25-MAR-2003 (revised)
DT 03-OCT-1995 (first entry)
XX
DE Human hek-L protein.
XX
KW Ligand; cell surface; tyrosine kinase receptor; tumorigenesis; immunogen.
XX
OS Homo sapiens.

XX Key Location/Qualifiers
FH Peptide 1..19
FT /note="signal peptide"
FT Protein 20..238
XX
XX NO506065-A1.
XX
XX PD 02-MAR-1995.
XX
XX PF 17-AUG-1994; 94MO-US009282.
XX
XX PR 20-AUG-1993; 93US-00109745.
XX PR 30-AUG-1993; 93US-00114426.
XX PR 03-DEC-1993; 93US-00161132.
XX PR 09-MAY-1994; 94US-00240124.
XX
XX PA (IMNV) IMMUNEX CORP.
XX
XX PI Beckmann MP, Cerretti DP;
XX
XX DR WPI, 1995-106811/14.
XX DR N-PSDB; AAO85887.
XX
XX PT New isolated DNA encoding hek-L protein or its fusion products - useful
PT as assay reagent or for carrying therapeutic and diagnostic compounds to
PT leukaemia cells.
XX
XX PS Claim 21; Page 36; 45pp; English.
XX
XX CC The sequence is that of a novel protein designated hek-L, a protein that
CC can bind hek (a cell surface receptor tyrosine kinase). Hek-L is the
CC first known ligand for hek and can be used to study cellular processes
CC regulated by hek (which may be involved in tumorigenesis). It is also an
CC immunogen for antibody production, as a reagent for detecting hek or hek-
CC L in vitro assays, to determine binding of hek proteins, to purify hek
CC proteins, and to carry diagnostic or cytotoxic agents to particular
CC leukaemia cells that express the hek antigen. Hek-L also binds the ek
CC tyrosine kinase receptors. See also AAR71482. (Updated on 25-MAR-2003 to
CC correct PN field.)
XX
XX SQ Sequence 238 AA;

Query Match 100.0%; Score 1207; DB 2; Length 238;
Best Local Similarity 100.0%; Pred. No. 1.8e-115;
Matches 219; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	MAAAPLLLLLVLPVPLPLPILAAQPGGALGNRAVYVNSNGHILREGYTVQVNVNDYLD	60
DB	1	MAAAPLLLLLVLPVPLPLPILAAQPGGALGNRAVYVNSNGHILREGYTVQVNVNDYLD	60
QY	61	IYCPHYNSGCVGAGPGGAGGAEQVYLVMVSNRGYRTCAAGCFRWECDNRDPAHSP	120
DB	61	IYCPHYNSGCVGAGPGGAGGAEQVYLVMVSNRGYRTCAAGCFRWECDNRDPAHSP	120
QY	121	KSEKFORISAEISLGYEFPAAGHEYYIISTPTNHLAKCLRMKYPVCCASHSNGEVPPT	180
DB	121	KSEKFORISAEISLGYEFPAAGHEYYIISTPTNHLAKCLRMKYPVCCASHSNGEVPPT	180
QY	181	LPQFTMGPNVKINVLSDPFGENPQVKLEKISIGTSFKR	219
DB	181	LPQFTMGPNVKINVLSDPFGENPQVKLEKISIGTSFKR	219

RESULT 2
ID AAG79439 standard; protein; 238 AA.
XX
AC AAG79439;
XX
DT 25-OCT-2002 (first entry)
XX
DE Breast/colon cancer associated protein.

XX Breast; colon; cancer; diagnosis; colorectal; prognosis; gene therapy;
 KW CHA4; CBK8.
 XX Homo sapiens.
 OS
 XX
 XX Key Location/Qualifiers
 FT Misc-difference 6..238
 PN /note="Encoded by ABA00091"
 WO200259609-A2.
 XX
 XX 01-AUG-2002.
 PD
 XX
 XX 10-DEC-2001; 2001WO-US048368.
 PF
 XX 08-DEC-2000; 2000US-00733756.
 PR 08-DEC-2000; 2000US-00733757.
 XX
 XX (EOSB-) EOS BIOTECHNOLOGY INC.
 PA
 XX Mack DH, Gish KC, Wilson KE;
 PI
 XX WPI; 2002-599815/64.
 DR
 XX N-PSDB; ABA00091.
 PT
 PT Diagnosing cancer by determining the expression profile gene that encodes
 PT the CHA4 or CBK8 protein, useful for the treatment and prognosis of
 PT breast and colorectal cancer.
 PS
 XX
 XX Disclosure; Fig 2; 95pp; English.
 XX
 XX The sequences given in ABA00086-92 represent sequences which have been
 CC associated with either breast and/or colon cancer. These sequences may be
 CC used in the method of the invention for diagnosing cancer. The method
 CC comprises determining the expression of one or more specified genes in a
 CC colorectal tissue sample, or in a breast tissue sample, of a first
 CC individual. The method of the invention is useful for the diagnosis,
 CC treatment and prognosis of breast and/or colorectal cancer. The invention
 CC also provides for methods useful for modulating activity and inhibiting
 CC breast and/or colorectal cancer. Prior methods of diagnosing and
 CC prognosticating breast and colorectal cancer have been limited and
 CC problematic. The present methods of gene therapy using novel sequences
 CC for the diagnosis and prognosis of cancer, including screening for drug
 CC candidates and bioactive agents that modulates CHA4 or CBK8, makes their
 CC use and purpose limitless and more specific especially in the area of
 CC grading prognostic factors. This sequence represents CHA4 which has the
 CC bioactivity of a breast cancer modulating protein
 CC
 CC Sequence 238 AA:
 SQ
 XX
 XX Query Match
 XX Best Local Similarity 100.0%; Score 1207; DB 5; Length 238;
 XX Matches 219; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAAAPLLILLVLPVPLPLAOGPGALGNRAVYVWNSNQHRLREGYTVQVAVNDYLD 60
 Db 1 MAAAPLLILLVLPVPLPLAOGPGALGNRAVYVWNSNQHRLREGYTVQVAVNDYLD 60
 QY 61 IYCPHYNSGVGPAGGPGGAGBOYVLYVWSRNGYRTCNASOGFKMECNRPDAPHSPI 120
 Db 61 IYCPHYNSGVGPAGGPGGAGBOYVLYVWSRNGYRTCNASOGFKMECNRPDAPHSPI 120
 QY 121 KFSEKFORYSASFSLGYEFHAGHEYYIISTPTNHLHWKCLRMKVFVCCASTSHSGEKVPVT 180
 Db 121 KFSEKFORYSASFSLGYEFHAGHEYYIISTPTNHLHWKCLRMKVFVCCASTSHSGEKVPVT 180
 QY 181 LPOFTMGPNVKINVLDEFGENPOVPLKESISGTSPEKR 219
 Db 181 LPOFTMGPNVKINVLDEFGENPOVPLKESISGTSPEKR 219

RESULT 3

ABP97190
 ID ABP97190 standard; protein; 238 AA.
 AC
 XX ABP97190;
 XX
 XX 01-JUL-2003 (first entry)
 DE
 XX Tumour-associated antigenic target protein TAT178 SEQ ID NO:72.
 KW Human; tumour-associated antigenic target; TAT; tumour; diagnosis;
 XX cancer.
 OS
 XX Homo sapiens.
 PA
 XX
 XX W02003024392-A2.
 PN
 XX
 XX 27-MAR-2003.
 PD
 XX
 XX 11-SEP-2002; 2002WO-US028859.
 PF
 XX
 XX 18-SEP-2001; 2001US-0323268P.
 PR 19-OCT-2001; 2001US-0339227P.
 PR 07-NOV-2001; 2001US-0336827P.
 PR 20-NOV-2001; 2001US-0331906P.
 PR 02-JAN-2002; 2002US-0345444P.
 PR 03-APR-2002; 2002US-0369724P.
 PR 19-AUG-2002; 2002US-0404809P.
 XX
 XX (GETH) GENENTECH INC.
 PA
 XX
 XX Frantz G, Hillan KT, Phillips HS, Polakis P, Spencer SD,
 PI Williams PM, Wu TD, Zhang Z;
 PI
 XX WPI; 2003-354551/33.
 DR
 XX N-PSDB; ACC49508.
 PT
 XX New antibodies against tumor-associated antigenic target polypeptide,
 PT useful for treating or diagnosing tumors or cancers in mammals, e.g.
 PT prostate cancer, lung cancer, prostate adenocarcinomas or renal cell
 PT carcinomas.
 PS
 XX Claim 2; Fig 72; 285pp; English.
 XX
 XX ACC49493 to ACC49552 encode the human tumour-associated antigenic target
 CC (TAT) proteins given in ABP97175 to ABP97234. The present invention
 CC describes an isolated antibody that binds to a polypeptide having at
 CC least 80 % sequence identity to any of the 60 150-800 residue amino acid
 CC sequences (S1), given in ABP97175 to ABP97234, comprising (S1), lacking
 CC its associated signal peptide, encoded by any of the 60 2000-3000 base
 CC cytosolic activity. The antibody can be used for treating or diagnosing
 CC tumours or cancers in mammals, e.g. prostate cancer, lung cancer, breast
 CC cancer, colon cancer, ovarian cancer, prostate adenocarcinomas, renal
 CC cell carcinomas, or thyroid cancer
 CC
 CC Sequence 238 AA:
 SQ
 XX
 XX Query Match
 XX Best Local Similarity 100.0%; Score 1207; DB 6; Length 238;
 XX Matches 219; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAAAPLLILLVLPVPLPLAOGPGALGNRAVYVWNSNQHRLREGYTVQVAVNDYLD 60
 Db 1 MAAAPLLILLVLPVPLPLAOGPGALGNRAVYVWNSNQHRLREGYTVQVAVNDYLD 60
 QY 61 IYCPHYNSGVGPAGGPGGAGBOYVLYVWSRNGYRTCNASOGFKMECNRPDAPHSPI 120
 Db 61 IYCPHYNSGVGPAGGPGGAGBOYVLYVWSRNGYRTCNASOGFKMECNRPDAPHSPI 120
 QY 121 KFSEKFORYSASFSLGYEFHAGHEYYIISTPTNHLHWKCLRMKVFVCCASTSHSGEKVPVT 180
 Db 121 KFSEKFORYSASFSLGYEFHAGHEYYIISTPTNHLHWKCLRMKVFVCCASTSHSGEKVPVT 180

QY 181 LPQFTMGPNVKINVLDEFEENPOVPLEKSIIGTSPKR 219
 DB 181 LPQFTMGPNVKINVLDEFEENPOVPLEKSIIGTSPKR 219
 RESULT 4
 ID ADN38812 standard; protein; 238 AA.
 AC ADN38812;
 XX
 DT 17-JUN-2004 (first entry)
 XX
 DE Cancer/angiogenesis/fibrosis-related polypeptide, SEQ ID NO:130.
 XX
 KW Human; differential expression; cancer; angiogenic disorder;
 KW fibrotic disorder; psoriasis; ischaemia; heart disease; atherosclerosis;
 KW inflammatory disease; autoimmune disease;
 KW retinal neovascularisation syndrome; scarring; uterine fibroid;
 KW detection; diagnosis; prognosis; drug screening; drug targeting;
 KW wound healing; contraception; cytostatic; cardiant; immunomodulatory;
 KW vulnery; gene therapy; vaccine.
 XX
 OS Homo sapiens.
 XX
 PN WO2003042661-A2.
 PD 22-MAY-2003.
 XX
 PF 13-NOV-2002; 2002WO-US036810.
 XX
 PR 13-NOV-2001; 2001US-0350666P.
 PR 21-NOV-2001; 2001US-0332464P.
 PR 29-NOV-2001; 2001US-0334393P.
 PR 03-DEC-2001; 2001US-0335394P.
 PR 14-DEC-2001; 2001US-0340376P.
 PR 08-JAN-2002; 2002US-0347211P.
 PR 10-JAN-2002; 2002US-0347349P.
 PR 08-FEB-2002; 2002US-0355250P.
 PR 13-FEB-2002; 2002US-0356714P.
 PR 20-FEB-2002; 2002US-0358077P.
 PR 29-MAR-2002; 2002US-0358809P.
 PR 04-APR-2002; 2002US-0370110P.
 PR 12-APR-2002; 2002US-0372246P.
 PR 05-JUN-2002; 2002US-0386614P.
 PR 16-JUL-2002; 2002US-0396839P.
 PR 22-JUL-2002; 2002US-0397775P.
 PR 22-JUL-2002; 2002US-0397845P.
 PR 09-SEP-2002; 2002US-0409450P.
 XX
 PA (EOSB-) EOS BIOTECHNOLOGY INC.
 XX
 PI Afar D, Aziz N, Ginsburg WM, Gish KC, Glyne R, Hevezi PA;
 PI Mack DH, Murray R, Watson SR, Wilson KE, Zlotnick A;
 XX
 DR WPI; 2003-468649/44.
 DR N-PSDB; ADN38811.
 XX
 PT Determining the presence or absence of a pathological cell in a patient,
 PT useful for diagnosing, prognosing or treating cancer, comprises detecting
 PT a nucleic acid in a biological sample.
 XX
 XX Claim 12; SEQ ID NO 130; 1385bp; English.
 XX
 CC The invention relates to nucleic acids and proteins (ADN38683-ADN40064)
 CC whose expression is upregulated or downregulated in specific cancers or
 CC other diseases such as angiogenic or fibrotic disorders, and to methods
 CC of determining the presence or absence of a pathological cell in a
 CC patient by detecting a nucleic acid at least 80% identical to those of
 CC the invention or by detecting a polypeptide of the invention. The
 CC invention also relates to expression vectors and host cells comprising a
 CC nucleic acid of the invention; antibodies which specifically bind a
 CC polypeptide of the invention; use of such antibodies for drug targeting;

CC and methods of screening for modulators of activity or expression of the
 CC polypeptides and nucleic acids. The nucleic acids, polypeptides,
 CC antibodies and methods are useful for diagnosing, prognosing and treating
 CC cancer and other conditions such as psoriasis, ischaemia, heart disease,
 CC atherosclerosis, inflammatory diseases, autoimmune diseases, retinal
 CC neovascularisation syndromes, scarring and uterine fibroids. They may
 CC also be useful in wound healing and in contraception. The present
 CC sequence represents a polypeptide of the invention.
 CC
 SO Sequence 238 AA:
 Query Match 100.0%; Score 1207; DB 7; Length 238;
 Best Local Similarity 100.0%; Pred. No. 1.8e-115;
 Matches 219; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAAPPLLLLLLVVPLPLPLLAQPGGALNRHAYVWVNSNOHRRGGTYQVWVNYLD 60
 DB 1 MAAPPLLLLLLVVPLPLPLLAQPGGALNRHAYVWVNSNOHRRGGTYQVWVNYLD 60
 QY 61 IYCPHYNSGVGPGAGPGGCAEQVLYVWSRNGYRTCNASOGFKMECNRPHAPHSPI 120
 DB 61 IYCPHYNSGVGPGAGPGGCAEQVLYVWSRNGYRTCNASOGFKMECNRPHAPHSPI 120
 QY 121 KFSEKFGRYSAFSLGYEFHAGHEYYIISTPTNHLAMKCLRMKVFVCCASTSHSGKEVPT 180
 DB 121 KFSEKFGRYSAFSLGYEFHAGHEYYIISTPTNHLAMKCLRMKVFVCCASTSHSGKEVPT 180
 QY 181 LPQFTMGPNVKINVLDEFEENPOVPLEKSIIGTSPKR 219
 DB 181 LPQFTMGPNVKINVLDEFEENPOVPLEKSIIGTSPKR 219
 RESULT 5
 ADQ21666
 ID ADQ21666 standard; protein; 238 AA.
 XX
 AC ADQ21666;
 XX
 DT 26-AUG-2004 (first entry)
 XX
 DE Human soft tissue sarcoma-upregulated; protein - SEQ ID 4486.
 XX
 KW soft tissue sarcoma; cytostatic; gene therapy; vaccine; screening; human.
 XX
 OS Homo sapiens.
 XX
 PN WO2004048938-A2.
 XX
 PD 10-JUN-2004.
 XX
 PF 26-NOV-2003; 2003WO-US038193.
 XX
 PR 26-NOV-2002; 2002US-0429739P.
 XX
 PA (PROT-) PROTEIN DESIGN LABS INC.
 XX
 PI Aziz N, Ginsburg WM, Zlotnick A;
 XX
 DR WPI; 2004-441208/41.
 DR
 PT Early detection of soft tissue sarcoma comprises determining expression
 PT of a gene in a first soft tissue sample and a normal soft tissue sample
 PT and comparing the gene expression, also useful in treating soft tissue
 PT sarcoma.
 XX
 XX Example 2; SEQ ID NO 4486; 210bp; English.
 XX
 CC The invention relates to a novel method for detecting soft tissue sarcoma
 CC which comprises obtaining a first soft tissue sample from an individual
 CC and a normal soft tissue sample from the same or different individual,
 CC determining the expression of a gene in both samples and comparing the
 CC expression of the gene in both soft tissue samples, where a higher level
 CC of protein expression in the first soft tissue sample indicates the

CC Presence of soft tissue sarcoma. The method of the invention has
 CC cytostatic applications and may be useful for detecting soft tissue
 CC sarcoma, possibly via gene therapy or vaccine production. The nucleic
 CC acid sequences may be useful in diagnostic and screening applications.
 CC The current sequence is that of a human soft tissue sarcoma-upregulated
 CC protein of the invention. The current sequence is not shown within the
 CC specification per se but was submitted in CD format by the inventor.
 XX Sequence 238 AA;

Query Match

Best Local Similarity 100.0%; Score 1207; DB 8; Length 238;
 Matches 219; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAAPLLILLVLPVPLPLLAQSPGALGNRHAYVYNNSSNOHLRREGYTVQVNVNDYLD 60
 DB 1 MAAPLLILLVLPVPLPLLAQSPGALGNRHAYVYNNSSNOHLRREGYTVQVNVNDYLD 60
 QY 61 IYCPHYNSGVPGGAGPDPGGAGAEQVLYVWSRNGYRTCNASOGFRKRCNRPAPHSPT 120
 DB 61 IYCPHYNSGVPGGAGPDPGGAGAEQVLYVWSRNGYRTCNASOGFRKRCNRPAPHSPT 120
 QY 121 KFESEKQRYSAFSLGYEFHAGHEYYISTPTNHLMKLCRMKVFVCCASTSHSGEKVPPT 180
 DB 121 KFESEKQRYSAFSLGYEFHAGHEYYISTPTNHLMKLCRMKVFVCCASTSHSGEKVPPT 180
 QY 181 LPOFTMGPNVKINVLDEPGENPOVPKLEKSIISGTSIPKR 219
 DB 181 LPOFTMGPNVKINVLDEPGENPOVPKLEKSIISGTSIPKR 219

RESULT 6

AAR82605
 ID AAR82605 standard; protein; 234 AA.
 XX AAR82605;

DT 16-MAY-1996 (first entry)

DE Eph transmembrane tyrosine kinase family ligand, Efl-2.

KW Efl-2; EHKL-1; Eph transmembrane tyrosine kinase family ligand;
 OS neurological disorder; identification; diagnosis.

XX Homo sapiens.

FT Key
 FT Peptide
 FT Location/Qualifiers
 FT 1..30
 FT /label= signal_peptide

FT Misc-difference 158
 FT /note= "residue borders main conserved regions"

FT Region
 FT 218..235
 FT /note= "carboxy terminal hydrophobic GPI-recognition
 tail"

XX W09527060-A2.

XX 12-OCT-1995.

XX 04-APR-1995; 95WO-US004208.

XX 04-APR-1994; 94US-00222075.

XX 12-APR-1994; 94US-00222075.

XX 01-SEP-1994; 94US-00222402.

XX 21-OCT-1994; 94US-00299567.

XX (REG-) REGENERON PHARM INC.

XX Davis S, Gale N, Aldrich TH, Maisompierre PC, Goldfarb M,
 XX Yancopoulos GD,
 XX WPI; 1995-358635/46.
 XX N-PSDB; AAT03883.

XX Ligands which bind Eph family receptors - used in the diagnosis of
 FT neurological disorders.
 PS Disclosure; Fig 2; 58pp; English.

XX Efl-2 (also known as EHKL-1) is an Eph transmembrane tyrosine kinase
 CC family ligand. It has homology with B61 (Efl-1) (see AAR82604). Efl-2
 CC ends in a C-terminal hydrophobic sequence that appears to be a
 CC recognition sequence allowing it to be GPI-linked and thus lacking in an
 CC intracellular domain. Efl-2 is useful for identifying other ligands for
 CC a differential function and/or influencing the phenotype, such as growth
 CC and/or proliferation, of receptor bearing cells. They may be used in the
 CC diagnosis, and treatment of neurological disorders

XX Sequence 234 AA;

Query Match

Best Local Similarity 97.1%; Score 1172; DB 2; Length 234;
 Matches 215; Conservative 0; Mismatches 0; Indels 4; Gaps 1;

QY 1 MAAPLLILLVLPVPLPLLAQSPGALGNRHAYVYNNSSNOHLRREGYTVQVNVNDYLD 60
 DB 1 MAAPLLILLVLPVPLPLLAQSPGALGNRHAYVYNNSSNOHLRREGYTVQVNVNDYLD 60
 QY 61 IYCPHYNSGVPGGAGPDPGGAGAEQVLYVWSRNGYRTCNASOGFRKRCNRPAPHSPT 120
 DB 61 IYCPHYNSGVPGGAGPDPGGAGAEQVLYVWSRNGYRTCNASOGFRKRCNRPAPHSPT 120
 QY 121 KFESEKQRYSAFSLGYEFHAGHEYYISTPTNHLMKLCRMKVFVCCASTSHSGEKVPPT 180
 DB 121 KFESEKQRYSAFSLGYEFHAGHEYYISTPTNHLMKLCRMKVFVCCASTSHSGEKVPPT 180
 QY 181 LPOFTMGPNVKINVLDEPGENPOVPKLEKSIISGTSIPKR 219
 DB 181 LPOFTMGPNVKINVLDEPGENPOVPKLEKSIISGTSIPKR 219

Search Completed: April 19, 2005, 23:13:33
 Job time: 59.9368 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 19, 2005, 09:30:04 ; Search time 391.37 Seconds
(without alignments)
8258.620 Million cell updates/sec

Title:	US-09-904-954-3_COPY_28_573
Perfect score:	546
Sequence:	1 ATGCGCTGCTGCCCTGCT.....GAGGGGGGACACTCCAGC 546

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 6

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000
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Post-processing: Minimum Match 89%
Maximum Match 100%
Listing first 1500 summaries

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2: genseeqn1980s:*
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4: genseeqn2000s:*
5: genseeqn2001as:*
6: genseeqn2001bs:*
7: genseeqn2002as:*
8: genseeqn2002bs:*
9: genseeqn2003as:*
10: genseeqn2003bs:*
11: genseeqn2003cs:*
12: genseeqn2004as:*
13: genseeqn2004bs:*

```

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	546	100.0	635	2	AA085888	AAG85888 Human hel
2	546	100.0	1181	8	AB234863	AB234863 Coding seq
3	544.4	99.7	606	6	ABV78135	ABV78135 Human epi
4	544.4	99.7	606	6	ABZ37111	ABZ37111 Human epi
5	544.4	99.7	606	6	ABX09954	ABX09954 Human epi
6	544.4	99.7	606	6	ABL91676	ABL91676 Human pol

ALIGNMENTS

RESULT 1	
AAQ85888	
ID	AAQ85888 standard; CDNA to mRNA; 636 BP.
XX	
AC	AAQ85888;
XX	
DI	25-MAR-2003 (revised)
DT	03-OCT-1995 (first entry)
XX	

DE		Human hek-L protein cDNA clone C6.
XX		
KM		Ligand, cell surface; tyrosine kinase receptor; tumorigenesis; immunogen;
XX		ss.
OS	Homo sapiens.	
XX		
PH	Key	Location/Qualifiers
FT	CDS	28..633
FT		/*tag= a
FT	sig_peptide	28..93
FT		/*tag= b
FT	mat_peptide	94..630
FT		/*tag= c
XX		
PN	WO9506065-A1.	
PD	02-MAR-1995.	
XX		
XX	17-AUG-1994;	94MO-US009282.
PF		
PR	20-AUG-1993;	93US-.00109745.
PR	30-AUG-1993;	93US-.00114426.
PR	03-DEC-1993;	93US-.00161132.
PR	09-MAY-1994;	94US-.00240124.
XX		
PA	(IMNV) IMMUNEX CORP.	
XX		
PI	Beckmann MP, Cerretti DP,	
DR	WPI, 1995-106811/14.	
DR	P-PSDB; AAR71482.	
XX		
PT	New isolated DNA encoding hek-L protein or its fusion products - useful	
PT	as assay reagent or for carrying therapeutic and diagnostic compounds to	
PT	leukaemia cells.	
XX		
PS	Claim 3; Page 37; 45pp; English.	
CC	The sequence is that of a clone encoding hek-L protein, a protein that	
CC	can bind hek (a cell surface receptor tyrosine kinase). Hek-L is the	
CC	first known ligand for hek and can be used to study cellular processes	
CC	regulated by hek (which may be involved in tumorigenesis). It is also an	
CC	immunogen for antibody production, as a reagent for detecting hek or hek-	
CC	L in vitro assays, to determine binding of hek proteins, to purify hek	
CC	proteins, and to carry diagnostic or cytotoxic agents to particular	
CC	leukemia cells that express the hek antigen. Hek-L also binds the elk	
CC	tyrosine kinase receptors. See also AA085887. (updated on 25-MAR-2003 to	
CC	correct FN field.)	
SQ	Sequence 636 BP; 102 A; 202 C; 186 G; 146 T; 0 U; 0 Other;	
Query Match	100.0%; Score 546; DB 2; Length 636;	
Best Local Similarity	100.0%; Pred. NO.2.1e-138; Mismatches 0; Gaps 0;	
Matches 546;	Conservative 0; Indels 0;	
1	ATGGGCGTGCAGCCCTTCGCACTGTCTTGAGCGCGGTTCTTCGACTCCCTCTG 60	
28	ATGGGCGTGCAGCCCTTCGCACTGTCTTCGAGACGTCTTCGCGCGGTTCTTCGCTCCCTCTG 87	
61	GCGCGGCGCTTCAGCCTTCGCCACGTAAGTCTTAGTGAATCTCAGTAACCCAGGTTGCT 120	
88	GCGCGGCGCTTCAGCCTTCGCCACGTAAGTCTTAGTGAATCTCAGTAACCCAGGTTGCT 147	
121	CGAAGAAGCCCGGTGTGAGTGTGGGCTCAAGATTAACTGAAGCATTTGTGCCCCCAC 180	
148	CGAAGAAGCCCGGTGTGAGTGTGGGCTCAAGATTAACTGAAGCATTTGTGCCCCCAC 207	
181	TACGAAGGCCCGAGGAGCCCTTGAAGGAGCCCGAGAGCGTTTGCTTTGTACATGTAAGACTG 240	
208	TACGAAGGCCCGAGGAGCCCTTGAAGGAGCCCGAGAGCGTTTGCTTTGTACATGTAAGACTG 267	
241	CGAGGCTATGAGTCTCTGCGAGGCAAGAGGACCCCGGCGCTAACAGCGCTGGGTGTCTCC 300	

Db 268 CAGGCTATGAGTCTCTCCAGAGAGAGGCCCCGCTTACAGCGCTGGGTGTCTCC 327
 QY 301 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAGATTCAGCGCTTACACCTTTCTCCCTC 327
 Db 328 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAGATTCAGCGCTTACACCTTTCTCCCTC 360
 QY 361 GGCCTTGAAGTCTTACCTGAGAGACTTCTACATCTCGGTGCCACTCCAGAGAGT 387
 Db 388 GGCCTTGAAGTCTTACCTGAGAGACTTCTACATCTCGGTGCCACTCCAGAGAGT 420
 QY 421 TCTGCCAGAGTCTTGAAGGCTCCAGGTGTCTGTCTGTCAGAGAGAGAGTCTGATCA 447
 Db 448 TCTGCCAGAGTCTTGAAGGCTCCAGGTGTCTGTCTGTCAGAGAGAGAGTCTGATCA 480
 QY 481 GCCCATCTCTGTTGGAGCCCTGAGAGAGTGGCACTCAGGTGGCGAGGGGAGACACT 507
 Db 508 GCCCATCTCTGTTGGAGCCCTGAGAGAGTGGCACTCAGGTGGCGAGGGGAGACACT 540
 QY 541 CCCAGC 546
 Db 568 CCCAGC 573

RESULT 2

AB234863
 ID AB234863 standard; cDNA; 1181 BP.
 AC AB234863;
 DT 04-FEB-2003 (first entry)

Coding sequence SEQ ID 221, differentially expressed in osteogenesis.

KW Osteoparctic; osteogenesis modulator; gene therapy; osteogenesis;
 KW osteoporosis; bone disease; upregulator; human; ephrin-ephrin; ss.
 OS Homo sapiens.
 PN W0200281745-A2.
 PD 17-OCT-2002.

PR 05-APR-2002; 2002WO-1B002211.
 PR 05-APR-2001; 2001US-0281400P.
 (AVER) AVENTIS PHARMA SA.

PI Garcia T, Roman Roman S, Baron R, Call K, Theilhaber J;
 PI Connolly T, Jackson A, Bushnell SE, Rawadi G;
 DR WPI; 2003-058567/05.

PT Novel isolated nucleic acid upregulated/downregulated in osteogenesis.
 PT useful for bone disease therapy in subject.
 PS Claim 1; Page 214; 237pp; English.

CC The present invention relates to novel nucleotide sequences, which are
 CC differentially expressed in models of osteogenesis upon being put in
 CC contact with a stimulator of osteogenesis. This sequence is one
 CC disease in a patient, promoting osteogenesis and/or preventing
 CC osteoporosis/bone disease. The present sequence encodes a Ephrin-ephrin
 CC family protein.

Sequence 1181 BP; 234 A; 353 C; 331 G; 263 T; 0 U; 0 Other;
 Query Match Best Local Similarity 100.0%; Score 546; DB 8; Length 1181;
 Matches 546; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCGGCTGCTGCCCCCTGCTGCGAAGTCTCTGAGGCGGGCTCTCGGCTCCCTCTG 60
 Db 28 ATGCGGCTGCTGCCCCCTGCTGCGAAGTCTCTGAGGCGGGCTCTCGGCTCCCTCTG 87
 QY 61 CGCGGGGGCTCCAGCTTCGCGCAGTACGTAAGTCTGAGTAAGTCCAGAGTGGTCT 120
 Db 88 CGCGGGGGCTCCAGCTTCGCGCAGTACGTAAGTCTGAGTAAGTCCAGAGTGGTCT 147
 QY 121 CGAGAGAGCGCGGTGAGGAGCTGGGCTCAAGATTAAGTCAATGCTGCCCCAC 180
 Db 148 CGAGAGAGCGCGGTGAGGAGCTGGGCTCAAGATTAAGTCAATGCTGCCCCAC 207
 QY 181 TACGAAGCGCCAGGGCCCCCTGAGGGCCCCGAGAGGTTTCTGTAATGAGTGA 240
 Db 208 TACGAAGCGCCAGGGCCCCCTGAGGGCCCCGAGAGGTTTCTGTAATGAGTGA 267
 QY 241 CCAAGCTATAGTCTGCGCAGGAGGCCCCCGGGCTTACAGCGCTGAGTGTCTCC 300
 Db 268 CCAAGCTATAGTCTGCGCAGGAGGCCCCCGGGCTTACAGCGCTGAGTGTCTCC 327
 QY 301 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAGATTCAGCGCTTACACCTTTCTCCCTC 360
 Db 328 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAGATTCAGCGCTTACACCTTTCTCCCTC 387
 QY 361 GGCCTTGAAGTCTTACCTGAGAGACTTCTACATCTCGGTGCCACTCCAGAGAGT 420
 Db 388 GGCCTTGAAGTCTTACCTGAGAGACTTCTACATCTCGGTGCCACTCCAGAGAGT 447
 QY 421 TCTGCCAGAGTCTTGAAGGCTCCAGGTGTCTGTCTGTCAGAGAGAGTCTGATCA 480
 Db 448 TCTGCCAGAGTCTTGAAGGCTCCAGGTGTCTGTCTGTCAGAGAGAGTCTGATCA 507
 QY 481 GCCCATCTCTGTTGGAGCCCTGAGAGAGTGGCACTCAGGTGGCGAGGGGAGACACT 540
 Db 508 GCCCATCTCTGTTGGAGCCCTGAGAGAGTGGCACTCAGGTGGCGAGGGGAGACACT 567
 QY 541 CCCAGC 546
 Db 568 CCCAGC 573

RESULT 3

ABV78135
 ID ABV78135 standard; DNA; 606 BP.
 AC ABV78135;
 DT 15-NOV-2002 (first entry)

DE Human ephrin-A3 DNA SEQ ID NO 19.
 KW RNA inhibition; dermal; gene expression inhibitor; oncogene; cytostatic;
 KW vitruclide; protozoaside; gene; ds.

OS Homo sapiens.
 PN W0200255693-A2.
 PD 18-JUL-2002.

PR 09-JAN-2002; 2002WO-EP000152.
 PR 26-OCT-2001; 2001DE-01000586.
 PR 29-NOV-2001; 2001DE-01055280.
 PR 07-DEC-2001; 2001DE-01058411.

PA (RIBO-) RIBOPHARMA AG.
 PI Kreutzler R, Limmer S, Rost S, Hadwiger P;
 DR WPI; 2002-590671/63.

PT Inhibiting expression of target gene, useful e.g. for inhibiting
PT oncogenes, by administering double-stranded RNA complementary to the
PT target and having an overhang.

PS Claim 10; Page 123; 203pp; German

The invention relates to inhibiting expression of a target gene (1) in cell by introducing an inhibitory RNA (dsRNA1) having a double-stranded structure of at most 49 consecutive bases. At least part of one strand (ss1) of dsRNA1 is complementary to (1) and at least one end of dsRNA1 has an overhang of 1-4 nucleotides. The method is used to inhibit the expression of a wide range of genes, e.g. oncogenes, cytokine genes etc. in humans, also genes in plasmidium or in viruses or plants that are pathogenic for humans, animals or plants. Introducing an overhang into dsRNA greatly increases effectiveness for inhibiting gene expression, both in vivo and in vitro and also increases stability and thus the effective concentration inside the cell. The present sequence is that of a gene related to the invention

Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match	99.78%	Score 544.4;	DB 6;	Length 606;
PostgreSQL Similarity:	00.00%	Word Weight 128		

Matches 545; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY	1	ATGGGGCTGCGACCCCTGCTGCGAGACTGTCCTCTGGGGCGGGTTCCTGGCTCCCTCTG	60
Db	1	ATGGGGCTGCGACCCCTGCTGCGAGACTGTCCTCTGGGGCGGGTTCCTGGCTCCCTCTG	60
QY	61	CGGGGGGCTTCAGCCTTCGCGCAGTACTTCTACTGGAACTCAGTAACCCAGATTGCTT	120
Db	61	CGGGGGGCTTCAGCCTTCGCGCAGTACTTCTACTGGAACTCAGTAACCCAGATTGCTT	120
QY	121	CGAGGAACCGCGGTGTGTGAGACTGGGCGCTCAACCAATTACCTAGACATTTGTCTGCCCCAC	180
Db	121	CGAGGAACCGCGGTGTGTGAGACTGGGCGCTCAACCAATTACCTAGACATTTGTCTGCCCCAC	180
QY	181	TACGAAAGCCACGAGGCGCCCTGAGAGGCCCGCAGACGTTTGCCTTTGTACATGGTGGACTGG	240
Db	181	TACGAAAGCCACGAGGCGCCCTGAGAGGCCCGCAGACGTTTGCCTTTGTACATGGTGGACTGG	240
QY	241	CCAGGCTATGAGTCTTGCCAGGCAAGAGGCCCGCGGCGCTTCAACAGCGCTGGGTGTCTCC	300
Db	241	CCAGGCTATGAGTCTTGCCAGGCAAGAGGCCCGCGGCGCTTCAACAGCGCTGGGTGTCTCC	300
QY	301	CTGGCCTTTGGCCATGTTCATTTCTCAGAGAAATTGAGGCGCTTACACCTTTCTCCCTC	360
Db	301	CTGGCCTTTGGCCATGTTCATTTCTCAGAGAAATTGAGGCGCTTACACCTTTCTCCCTC	360
QY	361	GGCTTTGAGTCTTACCTGAGAGACTTACTACTACATCTCGGTGCCACTCCAGAGAGT	420
Db	361	GGCTTTGAGTCTTACCTGAGAGACTTACTACTACTACATCTCGGTGCCACTCCAGAGAGT	420
QY	421	TCTGGCCAGTCTTGAAGGCTTCAGAGTGTCTGTCTGCTCAAGAGAGGAACTTGAATCA	480
Db	421	TCTGGCCAGTCTTGAAGGCTTCAGAGTGTCTGTCTGCTCAAGAGAGGAACTTGAATCA	480
QY	481	GCCCATCTCTTTGGGAGCCCTGAGAGAGAGTGGCATATAGAGGTGGCGAGGGGGGAGCACT	540
Db	481	GCCCATCTCTTTGGGAGCCCTGAGAGAGAGTGGCATATAGAGGTGGCGAGGGGGGAGCACT	540
QY	541	CCGAGC 546	
Db	541	CCGAGC 546	

RESULT 4

AB235711

ID AB235711 standard; DNA; 606 BP.

XX AB235711;

XX 07-FEB-2003 (first entry)

XX	Human ephrin A3 encoding polynucleotide SEQ ID NO 19.
DE	

KM Double stranded RNA; dsRNA; RNA inhibition; cytostatic; virucide;
KM protozooid; gene expression; antisense; tumour; infection; plasmodium
KM virus; anti-GFP; human; HIV; human immunodeficiency virus;
KM Hepatitis C virus; human papilloma virus; gene; ds.

OS Homo sapiens

PN DE10100588-A1.

PD 18-JUL-2002.

PF 09-JAN-2001; 2001DE-01000588.

PR 09-JAN-2001; 2001DE-01000588.

PA (RIBO-) RIBOPHARMA AG.

PI Kreutzer R, Limmer S, Rost S, Hadwiger P;

WPI; 2002-683450/74.

PT Inhibiting expression of target genes, useful e.g. for treating tumors,
PT by introducing into cells two double-stranded RNAs that are complementary
PT to the target.

PS Claim 13; Page 19-20; 100pp; German.

The invention relates to inhibiting expression of a target gene in a cell by introducing at least two oligonucleotides (dsRNA and II'), both with a double-stranded (ds) structure of at most 49 sequential nucleotides per strand. At least part of one strand (S1, S2) of the ds structures in each pair, at least part of one strand (S1, S2) of the ds structures in each pair, are complementary to regions in the target gene. The method uses antisense inhibition of gene expression using double stranded RNA inhibition (RNAi). The method is particularly useful to treat tumours or infections, especially by Paramodum or viruses/viroids (pathogenic organisms, animals or plants). The method provides more effective inhibition of expression than known methods using a single dsRNA, even at very low concentrations. When dsRNA has at least one unpaired nucleotide at the end, stability (and thus effective concentration in the cell) is improved and efficiency can be increased further by pretreating the cells with interferon. The present sequence is that of a target DNA of the invention

Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match 99.7%; Score 544.4; DB 6; Length 606;

Matches 545; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy	1	ATGGGCGTCTGCTCCCGCTGCGCGGAATGTCCTCTGGGGCGGGTTCCTCGAGCTCCCTCTG	60
Db	1	ATGGGCGTCTGCTCCCGCTGCGCGGAATGTCCTCTGGGGCGGGTTCCTCGAGCTCCCTCTG	60
Qy	61	CGCGGGGGCTCTCCAGCTCTCGCGCAGTAGTAGTCTACCTGGAATCCAGAGTGGCTT	120
Db	61	CGCGGGGGCTCTCCAGCTCTCGCGCAGTAGTAGTCTACCTGGAATCCAGAGTGGCTT	120
Qy	121	CGAGGAGACGCGCGTGTGGAGCTGGGCTCAACGATTACCTAGACATTGTCTGCCCCAC	180
Db	121	CGAGGAGACGCGCGTGTGGAGCTGGGCTCAACGATTACCTAGACATTGTCTGCCCCAC	180
Qy	181	TACGAAAGCCCAAGGGCCCCCTGAGAGGGCCCGAGACGTTGGTTGTACATAGTGGACATGG	240
Db	181	TACGAAAGCCCAAGGGCCCCCTGAGAGGGCCCGAGACGTTGGTTGTACATAGTGGACATGG	240
Qy	241	CCAAGCTATGAGTCTCTCCAGGACAGAGAGGCCCGGGCTTACAAGCGCTGGGTGTGCTCC	300
Db	241	CCAAGCTATGAGTCTCTCCAGGACAGAGAGGCCCGGGCTTACAAGCGCTGGGTGTGCTCC	300
Qy	301	CTGCGCTTTGGCCATGTTCAATTCTCAGAGAGATTCAAGCGCTTCAACCTTTTCTCCCTC	360

Db 301 CTGCCCTTGGCAGTTCATTTCTCAGAGAGATTGACGGCTTACACCTTCTCCCTC 360
 QY 361 GGCCTTGAGTTCTTACCTGAGAGACTTACTACTACTGCTGGCTCCCTCCAGAGAGT 420
 Db 361 GGCCTTGAGTTCTTACCTGAGAGACTTACTACTACTGCTGGCTCCCTCCAGAGAGT 420
 QY 421 TCTGGCCAGTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 480
 Db 421 TCTGGCCAGTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 480
 QY 481 GGCCTTGAGTTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 540
 Db 481 GGCCTTGAGTTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 540
 QY 541 CCCAGC 546
 Db 541 CCCAGC 546

RESULT 5
 ABX09954
 ID ABX09954 standard; DNA; 606 BP.
 AC ABX09954;
 XX
 DT 23-JAN-2003 (first entry)
 DE
 XX Human ephrin A3 DNA fragment SEQ ID 19.
 KM
 XX Oligoribonucleotide, interferon; oncogene; cytokine; id; developmental;
 XX prion; inhibition; human; ds.
 OS Homo sapiens.

XX DE10100587-C1.
 XX
 PN 21-NOV-2002.
 PD
 PF 09-JAN-2001; 2001DE-01000587.
 PR 09-JAN-2001; 2001DE-01000587.
 XX
 PA (RIBO-) RIBOPHARMA AG.
 XX
 PI Kreutzer R, Limmer S, Rost S, Hadwiger P,
 DR WPI; 2002-742209/81.

XX Inhibiting expression of target genes, e.g. oncogenes, in cells, by
 PT introduction of complementary double-stranded oligoribonucleotide, after
 PT treating the cell with interferon.
 PS Disclosure: Page 24-25; 98pp; German.

XX This invention describes a novel method for inhibiting expression of a
 CC target gene by introducing into the cell that contains the target gene at
 CC least one oligoribonucleotide (dsRNA) that has a double-stranded (ds)
 CC structure of not more than 49 consecutive nucleotides (nt), where at
 CC least a segment of one strand of the ds structure is complementary with
 CC the target gene and the cells are treated with interferon before
 CC introduction of dsRNA. The method is used to inhibit expression of
 CC target genes, particularly oncogenes, cytokine genes, id (not defined)
 CC protein genes, developmental or prion genes, or genes expressed in
 CC pathogenic organisms (particularly plasmids) or in viruses or viroids
 CC (pathogenic in humans, animals or plants). Treating the cells with
 CC interferon greatly increases the extent to which dsRNA can inhibit
 CC expression of the target genes, and the effect is even greater when dsRNA
 CC are modified to increase their stability. ABX09954-ABX10075 represent
 CC gene fragments used to illustrate the method of the invention
 SQ Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match

99.7%; Score 544.4; DB 6; Length 606;

Best Local Similarity 99.8%; Pred. No. 5,5e-138;
 Matches 545; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCGGCTGCTGCGCCCGCTGCGGAGACTGCTCTTGTGGCCCGCTTCTCCCTCTG 60
 Db 1 ATGCGGCTGCTGCGCCCGCTGCGGAGACTGCTCTTGTGGCCCGCTTCTCCCTCTG 60
 QY 61 CGCGGGGGGCTCCAGGCTCTCGCCAGTGTACTAGTCAATCCAGTACCCAGAGTGTCT 120
 Db 61 CGCGGGGGGCTCCAGGCTCTCGCCAGTGTACTAGTCAATCCAGTACCCAGAGTGTCT 120
 QY 121 CGAGAGAGACCGCTGTGTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTG 180
 Db 121 CGAGAGAGACCGCTGTGTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTGAGCTG 180
 QY 181 TACGAGAGCCCAAGGCGCCCTGAGGGCCCGGAGAGCTTGTGTTGTTGTTGTTGTTG 240
 Db 181 TACGAGAGCCCAAGGCGCCCTGAGGGCCCGGAGAGCTTGTGTTGTTGTTGTTGTTG 240
 QY 241 CCAGGCTATAGTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 300
 Db 241 CCAGGCTATAGTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 300
 QY 301 CTGCGCTTGTGGCCATGTTCAATTCTCAGAGAGAAAGTTCAGCGCTTCTCCCTC 360
 Db 301 CTGCGCTTGTGGCCATGTTCAATTCTCAGAGAGAAAGTTCAGCGCTTCTCCCTC 360
 QY 361 GGCCTTGAGTTCTTACCTGAGAGACTTACTACTACTGCTGGCTCCCTCCAGAGAGT 420
 Db 361 GGCCTTGAGTTCTTACCTGAGAGACTTACTACTACTGCTGGCTCCCTCCAGAGAGT 420
 QY 421 TCTGGCCAGTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 480
 Db 421 TCTGGCCAGTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 480
 QY 481 GGCCTTGAGTTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 540
 Db 481 GGCCTTGAGTTCTTGAAGGCTCCAGGCTGTCTGCTGCTGCTGCAAGAGAGAAAGTCTAGTCA 540
 QY 541 CCCAGC 546
 Db 541 CCCAGC 546

RESULT 6

ABL91676
 ID ABL91676 standard; DNA; 606 BP.
 AC ABL91676;
 XX
 DT 28-MAY-2002 (first entry)
 DE
 XX Human polynucleotide SEQ ID NO 19.
 KM Human; HIV; HCV; gene expression; oligoribonucleotide; tumour; pathogen;
 KM Plasmidum; virus; viroid; cytokine; prion; antisense oligonucleotide;
 KM Cytostatic; virulence; protozoicide; antibacterial; ds.
 OS Homo sapiens.

XX DE10100586-C1.
 XX
 PN 11-APR-2002.
 PD
 PF 09-JAN-2001; 2001DE-01000586.
 PR 09-JAN-2001; 2001DE-01000586.
 XX
 PA (RIBO-) RIBOPHARMA AG.
 XX
 PI Kreutzer R, Limmer S, Rost S, Hadwiger P,
 DR WPI; 2002-270454/32.

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; ZIP: 98101

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COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/240,124
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/161,132
FILING DATE: 03-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
TELEX: 7556822
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 636 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHEetical: NO
ANTI-SENSE: NO
IMMEDIATE SOURCE:
CLONE: hek-L C6
FEATURES:
NAME/KEY: mat_peptide
LOCATION: 94..630
FEATURE:
NAME/KEY: CDS
LOCATION: 28..633
FEATURES:
NAME/KEY: sig_peptide
LOCATION: 28..93
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US-08-240-124-3

Query Match          100.0%; Score 546; DB 1; Length 636;
Best Local Similarity 100.0%; Pred. No. 1,9e-142;
Matches 546; Conservativity 0; Mismatches 0; Indels 0; Gaps 0;

    1 ATGGGCGTGCAGCCCTGTGTGGGAGACTGCTCTGGGGCGGGTTCTTGAGCCTCCCTCTG 60
      |||||
    28 ATGGGCGTGCAGCCCTGTGTGGGAGACTGCTCTGGGGCGGGTTCTTGAGCCTCCCTCTG 87
      |||||
    61 CGCGGGGCGTTCACAGCTTCGCGCACGTAAGTCTAATGAACTCAAGTAACCCGAGTTGCTT 120
      |||||
    88 CGCGGGGCGTTCACAGCTTCGCGCACGTAAGTCTAATGAACTCAAGTAACCCGAGTTGCTT 147
      |||||
    121 CGAGGAGACGCCGTGTGTGAAGCTGAGGCTCAACGATTACTTAGACATTGTCTGCCCCAAC 180
      |||||
    148 CGAGGAGACGCCGTGTGTGAAGCTGAGGCTCAACGATTACTTAGACATTGTCTGCCCCAAC 207
      |||||
    181 TAGCAGAGCCCGAGGGCCCCCGTAGAGGGCCCCGAGACGTTTGCTTTGATCATGTGTGACTGG 240
      |||||
    208 TACAGAGCCCGAGGGCCCCCGTAGAGGGCCCCGAGACGTTTGCTTTGATCATGTGTGACTGG 267
      |||||
    241 CCAGGCTATGAGTCTGTGCAGGACAGAGGCCCCCGGGCTTACAAGCGCTGGGTGTGCTCC 300
      |||||
    268 CCAAGGCTATGAGTCTGTGCAGGACAGAGGCCCCCGGGCTTACAAGCGCTGGGTGTGCTCC 327
      |||||
    301 CTGCGCTTTGGCCATGTTCATTCTCAGAGAAGATTACGGCTTCACACCTTTCTCCCTC 360

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RESULT 2
 US-08-453-943-3
 ; Sequence 3, Application US/08453943
 ; Patent No. 5738844
 GENERAL INFORMATION:
 APPLICANT: BECKMANN M. P.
 APPLICANT: CERRETTI, DOUGLAS P.
 TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
 NUMBER OF INVENTION: RECEPTOR HER
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: IMMUNEX CORPORATION
 STREET: 51 UNIVERSITY STREET
 CITY: SEATTLE
 STATE: WASHINGTON
 COUNTRY: USA
 ZIP: 98101
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: Apple Macintosh
 OPERATING SYSTEM: Apple System 7.1
 SOFTWARE: Microsoft Word for Apple, Version 5.1a
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/453,943
 FILING DATE: 30-MAY-1995
 CLASSIFICATION: 530
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/240,124
 FILING DATE: 09-MAY-1994
 APPLICATION NUMBER: US 08/161,132
 FILING DATE: 03-DEC-1993
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/114,426
 FILING DATE: 30-AUG-1993
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/109,745
 FILING DATE: 20-AUG-1993
 ATTORNEY/AGENT INFORMATION:
 NAME: SEESE, KATHRYN A.
 REGISTRATION NUMBER: 32,172
 REFERENCE/DOCKET NUMBER: 2814-C
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (206) 587-0030
 TELEFAX: (206) 233-0644
 TELEX: 756822
 INFORMATION FOR SEQ ID NO: 3:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 636 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: cDNA to mRNA
 HYPOTHEICAL: NO

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? ? ANTI-SENSE: NO
? ? IMMEDIATE SOURCE:
? ? CLONE: hek-L C6
? ? FEATURE:
? ? NAME/KEY: mat_peptide
? ? LOCATION: 94..630
? ? FEATURE:
? ? NAME/KEY: CDS
? ? LOCATION: 28..633
? ? FEATURE:
? ? NAME/KEY: sig_peptide
? ? LOCATION: 28..93
US-08-453-943-3

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Query Match      100.0%; Score 546; DB 1; Length 636;
Best Local Similarity 100.0%; Pred. No. 1.9e-142;
Matches 546; Conservative 0.

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[illegible]

RESULT 3
US-09-057-121-3
Sequence 3, Application US/09057121
Patent No. 5693110
GENERAL INFORMATION:
APPLICANT: BECKMANN, M. P.
APPLICANT: CERRETTI, DOUGLAS P.
TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
NUMBER OF INVENTOR: RECEPTOR HEK
CORRESPONDENCE ADDRESSES: 4
ADDRESSEE: IMMUNEX CORPORATION
STREET: 51 UNIVERSITY STREET
CITY: SEATTLE

STATE: WASHINGTON
COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/057,121
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/240,124
FILING DATE:
APPLICATION NUMBER: US 08/161,132
FILING DATE: 03-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
TELEX: 756822
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 636 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
IMMEDIATE SOURCE:
CLONE: hek-L C6
FEATURE:
NAME/KEY: mat_peptide
LOCATION: 94..630
FEATURE:
NAME/KEY: CDS
LOCATION: 28..633
FEATURE:
NAME/KEY: sig_peptide
LOCATION: 28..93
US-09-057-121-3

Query Match 100.0%; Score 546; DB 2; Length 636;
Best Local Similarity 100.0%; Pred. No. 1.9e-142;
Matches 546; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCGGCTGCTGCGCCCTGCTGCGGACGCTGCTCTGCGGCGCGGCTTCTCGGCTCCCTCTG 60
DB 28 ATGCGGCTGCTGCGCCCTGCTGCGGACGCTGCTCTGCGGCGCGGCTTCTCGGCTCCCTCTG 87
QY 61 CGCGGGGGGCTTCAGGCTTCGCGGACGTAAGTCTAGTAAGTTCAGTAACCCAGGTTGCTT 120
DB 88 CGCGGGGGGCTTCAGGCTTCGCGGACGTAAGTCTAGTAAGTTCAGTAACCCAGGTTGCTT 147
QY 121 CGAGGAGACGCGCGGTGAGAGCTGAGGCTGAGGCTCAACGATTACCTAGACATTGCTGCCCCAC 180
DB 148 CGAGGAGACGCGCGGTGAGAGCTGAGGCTGAGGCTCAACGATTACCTAGACATTGCTGCCCCAC 207
QY 181 TACGAGGCCCGAGGCCCTGAGAGGCCCGAGACGTTTGTTGTAATGAGTGAAGTGG 240
DB 208 TACGAGGCCCGAGGCCCTGAGAGGCCCGAGACGTTTGTTGTAATGAGTGAAGTGG 267

QY 241 CCAGCTATGATGCTCTGCCAGGAGGCCCCCGGCTTACAGGCTGGGTGCTCC 300
DB 268 CCAGGCTATGATGCTCTGCCAGGAGGCCCCCGGCTTACAGGCTGGGTGCTCC 327
QY 301 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAAATTGAGGCTTTCACACTTTCTCCCTC 360
DB 328 CTGCCCTTTGGCCATGTTCAATTCTCAGAGAAATTGAGGCTTTCACACTTTCTCCCTC 387
QY 361 GCGTTGAGTCTTACTGAGAGACTTCTACTACATCGGTTGCCACTCCAGAGAT 420
DB 388 GCGTTGAGTCTTACTGAGAGACTTCTACTACATCGGTTGCCACTCCAGAGAT 447
QY 421 TCTGCCAGAGCTTGAAGGCTCCAGGTGTCTGTCTGTCGCAAGAGAGAGAGTGA 480
DB 448 TCTGCCAGAGCTTGAAGGCTCCAGGTGTCTGTCTGTCGCAAGAGAGAGAGTGA 507
QY 481 GCCATCTCTGTTGGAGCCCTGGAGAGAGTGCACATCAGGGTGGGAGGGGAGACT 540
DB 508 GCCATCTCTGTTGGAGCCCTGGAGAGAGTGCACATCAGGGTGGGAGGGGAGACT 567
QY 541 CCCAGC 546
DB 568 CCCAGC 573

RESULT 4
US-09-358-734-3
Sequence 3, Application US/09358734
Patent No. 6274117
GENERAL INFORMATION:
APPLICANT: BECKMANN, M. P.
TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
TITLE OF INVENTION: RECEPTOR HEK
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: IMMUNEX CORPORATION
STREET: 51 UNIVERSITY STREET
CITY: SEATTLE
STATE: WASHINGTON
COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/358,734
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/240,124
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
TELEX: 756822
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 636 base pairs
TYPE: nucleic acid
STRANDEDNESS: single

28..93

[illegible]

61 CCGGGGAGCTCGAGCTCCCGGCGCAGTAGTCTACTGAACTCCAGTAAACCCAGAGTTGCT 60
 88 CGCGGGAGCTCGAGCCCTCGGCGCAGTAGTCTACTGAACTCCAGTAAACCCAGAGTTGCT 120
 121 CGAGGAGACGCGGTGTGTGGAGCTGTGGAGCTTCAACGATTAAGTGAATGTCTGGCCCCAC 147
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 508 GCCATCTCTTGTGGAGCCCTGAGAGAGAGTGTGACATCAAGAGAGAGAGAGTCTGAGTCA 540
 541 CCTGAGC 546
 568 CCTGAGC 573

RESULT 5
 US-09-949-016-1291
 Sequence 1291, Application US/09949016
 Patent No. 681238
 GENERAL INFORMATION:
 APPLICANT: VENTER, J. Craig et al.
 TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 FILE REFERENCE: CLO01307
 CURRENT APPLICATION NUMBER: US/09/949,016
 CURRENT FILING DATE: 2000-04-14

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? PRIOR APPLICATION NUMBER: 60/241,755
? PRIOR FILING DATE: 2000-10-20
? PRIOR APPLICATION NUMBER: 60/237,768
? PRIOR FILING DATE: 2000-10-03
? PRIOR FILING DATE NUMBER: 60/231,498
? NUMBER OF SEQ ID NOS: 207012
? SOFTWARE: Paq ID NOS: 207012
? SEQ ID NO 1291
? LENGTH: 1182
? TYPE: DNA
? ORGANISM: Human
US-09-949-016-1291

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Best Local Similarity	99.8%	Pred. No. 6.6e-142;	
Matches 545; Conservative	0; Mismatches		

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26	ATGCGACTGCTGCCCTCTGCTGCGGACTGCTCTCGGCGCGGTTCCCTCGGCTCCCTCTCG	60
61	CGCGGGGCTTCAAGCCCTCGGCGGCACTGATCTACTGGAATCCGCTGAACCCGAGTTGGTT	87
88	CGCGGGGCTTCAAGCCCTCGGCGGCACTGATCTACTGGAATCCGCTGAACCCGAGTTGGTT	120
121	CGAGGAGACGCGGTTGGTGAAGCTGGGCTCAAGATTACTTAAGACATTGTCGCCCCC	147
148	CGAGGAGACGCGGTTGGTGAAGCTGGGCTCAAGATTACTTAAGACATTGTCGCCCCC	180
181	TACGAAAGCCGAGGCCCCCTGAGGGCCCCGAGAGCTTGTGTGAACAGTGTCTGCCCCC	207
208	TACGAAAGCCGAGGCCCCCTGAGGGCCCCGAGAGCTTGTGTGAACAGTGTCTGCCCCC	240
241	CGAGGCTATGATCTGTCGAGGAGAGGAGGCCCCGAGGCTTGTGTGAACAGTGTCTGCCCCC	267
268	CGAGGCTATGATCTGTCGAGGAGAGGAGGCCCCGAGGCTTGTGTGAACAGTGTCTGCCCCC	300
301	CTGCGCTTTGGCCATGTTCAATTCTCAAGAGATTCAGCGCTTACACTTTCTCCCTC	327
328	CTGCGCTTTGGCCATGTTCAATTCTCAAGAGATTCAGCGCTTACACTTTCTCCCTC	360
361	GGCTTTGAGTTCTTACCTGGAGAGACTTACTTCAATCTGGTGGCCACTCCAGAGAGT	387
388	GGCTTTGAGTTCTTACCTGGAGAGACTTACTTCAATCTGGTGGCCACTCCAGAGAGT	420
421	TTGGGCGAGGCTTGAAGGCTTCCAGGTGTCTGTCTGGCTGGCAAGAGAGAAAGTCTGAAGT	447
448	TTGGGCGAGGCTTGAAGGCTTCCAGGTGTCTGTCTGGCTGGCAAGAGAGAAAGTCTGAAGT	480
481	GCCCATCTGTTGGAGGCTTGAGAGAGAGGCAATCAGAGGTGGCGAGGGGGGAGACT	507
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568	CCGAGC 573	567

Search completed: April 19, 2005, 22:12:42
Job time : 118.434 secs


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ORIGIN
Query: 1

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Conservative	0		Indels	0
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DB	103	CGCGGGGGCTCGAGCTCCGCCGAGTGTACTGTGAATCTCAGTAACCCCGAGGTGTT	120	
QY	121	CGAGGAGACGCGCGGTGTGTAGTGTGGCTTCAGACATTTACTAGACATTTGTCTGCCCCAC	162	
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QY	181	TACGAAAGCCGAGGGCCCCCTGAGGCGCCCGACGCTTTGTCTTTCACGTGTGTGACCTCG	222	
DB	223	TACGAAAGCCGAGGGCCCCCTGAGGCGCCCGACGCTTTGTCTTTCACGTGTGTGACCTCG	222	
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DB	283	CCAGGCTATAGTCTGTCCAGCAGAGAGGCCCCCGGCGCTACAGGCTGTGTGTCTTC	240	
QY	301	CTGCGCTTTGGCATATGTTCAATCTGAGAGAGATTGAGCGCTTCACACCTTTCTCCCTC	342	
DB	343	CTGCGCTTTGGCATATGTTCAATCTGAGAGAGATTGAGCGCTTCACACCTTTCTCCCTC	342	
QY	361	GGCTTTAGTCTTAACTGTGAGAGCTTAACTAACTGTGAGAGCTTAACTGTGAGAGCTTAACT	402	
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RESULT 3	
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LOCUS	
DEFINITION	BQ928753
ACCESSION	AGNCOCURT_10036172 NIH_MGC_40 Homo sapiens mRNA
VERSION	5, mRNA sequence.
KEYWORDS	BQ928753
SOURCE	BQ928753.1 GI:22343784
ORGANISM	Homo sapiens (human)
REFERENCE	1 Homo sapiens
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
JOURNAL	NIH-MGC http://mgc.nci.nih.gov/
COMMENT	National Institutes of Health, Mammalian Gene Collection Project Contact: Robert

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        Location/Qualifiers
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            directionally, cDNA made by oligo-dT priming.
            Lining Hong in the laboratory of GSCACGAG(Ge). Library constructed by
            (Stratagene, Berkeley) using Zald M. Rubin (University
            Note: this is a NIH MGC Library." (Life
            Technologies).

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      99.7%; Score 544.4; Db 5; Length 965;
Best Local Similarity 99.8%; Pred. No. 1.4e-129;
Matches 545; Conservative 0; Mismatches 1; Indels 0; Gaps
QY      1 ATCGGCTGCTGCTCCCTCTGCGAGCTGTCCTTGGCGCGGTTCCCGGCTCCCTCTG
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QY      61 CGCGGGGCTTCAAGCTCCGCGCAGTATCTACTGGAATCCAGTAACCCAGTTGCTT
Db      116 CGCGGGGCTTCAAGCTCCGCGCAGTATCTACTGGAATCCAGTAACCCAGTTGCTT
QY      121 CAGAGAGACCGCTGTGTGAAGCTGGGCTCAAGGATTTACGTAGCATTTGCTGCTCCCGAC
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Db 176 CGAGGAGACGCCCTGTGAGCTGGGCTCAACGATTACTAGACATTTGCTGCCCCAC 235
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 Db 356 CTGCGCTTTGGCCATGTTCAATTTCTCAGAGAAATTACGCGCTTACACCTTTCTCC 415
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 Db 416 GGGTTGAGTTCTTACCTGAGAGACTTACTACATCTCGGTGCCACTCCAGAGAGT 475
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 Db 476 TCTGGCAGTCTTGAAGGCTCCAGTGTCTGTCTGTCAGAGAGAGAACTGAGTCA 535
 QY 481 GCCCATCTCTTTGGAGCCCTGAGAGAGAGTGCACATCAGGGTGGCGAGGGGGAGACT 540
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 QY 541 CCCAGC 546
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RESULT 4
 LOCUS CA454936 898 bp mRNA linear EST 12-NOV-2002
 DEFINITION AGENCOURT 10714679 MAPCL Homo sapiens cDNA clone IMAGE:6722424 5',
 CA454936
 mRNA sequence.
 ACCESSION CA454936
 VERSION CA454936.1 GI:24905152
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 898)
 NIH-MGC http://mgc.nci.nih.gov/
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgabs-remail.nih.gov
 Tissue Procurement: Kristi A. Eglund, Ira Pastan
 cDNA Library Preparation: Invitrogen Corp
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LLAM14285 row: b column: 24
 High quality sequence stop: 473.
 Location/Qualifiers
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 Directionally cloned. Priming method: oligo-dT. Average
 insert size: 1800 bp. Library amplification: 26,000 fold.
 Kristi A. Eglund, James J. Vincent, Robert Strausberg,

FEATURES
 source

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 COMMENT

ORIGIN
 Bungkok Lee & Ira Pastan: Discovery of new breast
 cancer genes encoding membrane and secreted proteins.
 Manuscript submitted."

Query Match 98.4%; Score 537; DB 6; Length 898;
 Best Local Similarity 98.9%; Pred. No. 1, 1e-127;
 Matches 540; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 ATGCGGCTGCTGCCCCCTGCGGAGACTGCTCTCGGGCGCGCTTCCCTGCTCCCTG 60
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 QY 181 TACGAAGGCCAGAGGCCCTCTGAGGGCCCGAGACGTTTGTATCATGTGACTGG 240
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 Db 296 CCAGGCTATGAGTCTGCGCAGGAGAGAGGCCCGGCTTCAAGCGCTGGTGTCTCC 355
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 Db 356 CTGCGCTTTGGCCATGTTCAATTTCTCAGAGAAATTACGCGCTTACACCTTTCTCC 415
 QY 361 GGGTTGAGTTCTTACCTGAGAGACTTACTACATCTCGGTGCCACTCCAGAGAGT 420
 Db 416 GGGTTGAGTTCTTACCTGAGAGACTTACTACATCTCGGTGCCACTCCAGAGAGT 475
 QY 421 TCTGGCAGTCTTGAAGGCTCCAGTGTCTGTCTGTCAGAGAGAGAACTGAGTCA 480
 Db 476 TCTGGCAGTCTTGAAGGCTCCAGTGTCTGTCTGTCAGAGAGAGAACTGAGTCA 535
 QY 481 GCCCATCTCTTTGGAGCCCTGAGAGAGAGTGCACATCAGGGTGGCGAGGGGGAGACT 540
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 BE883793
 mRNA sequence.
 ACCESSION BE883793
 VERSION BE883793.1 GI:10332569
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 707)
 NIH-MGC http://mgc.nci.nih.gov/
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgabs-remail.nih.gov
 Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be

Found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: L1AM9718 row: b column: 23
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 Location/Qualifiers
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 Best Local Similarity 99.6%; Pred. No. 1.6e-126;
 Matches 544; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

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 49 ATGCGCTGCTGCCCCCTGCGGAGCTGCTCTGGGCGGCGTTCGGCTCCCTCTG 60
 61 CGCGGGGCTCCAGGCTCGCCAGCTGCTCTGAGAGCTCCGAGCTCCCTCTG 108
 109 CGCGGGGCTCCAGGCTCGCCAGCTGCTCTGAGAGCTCCGAGCTCCCTCTG 108
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 VERSION B168252.1 GI:16041925
 KEYWORDS
 EST. Homo sapiens (human)
 SOURCE
 ORGANISM Homo sapiens

REFERENCE
 AUTHORS
 TITLE
 JOURNAL
 COMMENT
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 NIH-MGC <http://mgi.nci.nih.gov/>.
 National Institutes of Health, Mammalian Gene Collection (MGC)
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: sgabbs-remail.nih.gov
 Tissue Procurement: ATCC
 CDNA Library Preparation: Life Technologies, Inc.
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: L1AM1205 row: j column: 01
 High quality sequence stop: 806.
 Location/Qualifiers
 1..912

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 Note: this is a NIH_MGC Library."

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QY	541	CCCAGC	546
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DEFINITION	601431292F1 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:3916487 5', mRNA sequence.				

VERSION	BE890843.1	GI:10349570
KEYWORDS	EST.	
SOURCE	Homo sapiens	(human)
ORGANISM	Homo sapiens	

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 838)
NIH-MGC <http://mgc.nci.nih.gov/>.
National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999)
Contact: Robert Strausberg, Ph.D.

Email: CGAPds-r@mail.nih.gov
 Tissue Procurement: ATCC/DCTD/DNP
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: [The I.M.A.G.E. Consortium \(LNLN\)](http://The I.M.A.G.E. Consortium (LNLN))
 DNA Sequencing by: Invitrogen, Inc.
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNLN at:
<http://image.llnl.gov>
 Plate: LIM5741 row: h column: 24
 High quality sequence stop: 707.
 Location/Qualifiers

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                Technologies."

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Best Local Similarity	99.5% Pred. No. 1,2e-123;
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Db	71	ATGCGGCTGCTGCCTCCCTGCTGCAGACTGTCTCTGGAGCGCGTTCCTCGGCTCCCTCTG	13
Qy	61	CGGGGGGGCTTCAGCTCTCGGCACGTAGTCTACTGGAATCCAGTAACCCCAAGTTGCTT	12
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Qy	121	CGAGAGACGCGCTGTGTGAGCTGGGCGTCAACGATTACTAGACATTTGTCCGCCAC	18
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Qy	301	CTGCGCTTTGGCAATGTTCAATTCTAGAGAGATTCAGGCTTCACACCTTTCCTCTC	36

Db	370	CTGCGCTTTGGCGAATGTTCAATTCACAGAAAGATAGCGCTTCACACCCTTCTCCCTC	429
Qy	361	GGCTTTGAATTCCTTACCTGGAGAGACTTACTACTACATCTCTGGTGGCCACTCCAGAGAGT	420
Db	430	GGTTTGAATTCCTTACCTGGAGAGACTTACTACTACATCTCTGGTGGCCACTCCAGAGAGT	489
Qy	421	TCGGCGCATGCTTGAAGGCTCCAGGTGTCTGTCTCTGCTGCAGAGAGAGGAAAGTCTGAATCA	480
Db	490	TCGGCGCATGCTTGAAGGCTCCAGGTGTCTGTCTCTGCTGCAGAGAGAGGAAAGTCTGAATCA	549
Qy	481	GCCCATCTCTGTTGGAGGCCCTGGAGAGAGTGGACATCAGGCTGGCGA-GGGGGGGAAC	539
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Qy	540	TCCGAGC 546	
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DEFINITION	17000424723908 GRN_ES Homo sapiens				
ACCESSION	CN289864				
VERSION	CN289864.1	GI:47306278			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 663)
REFERENCE
AUTHORS Brandenberger R., Wei H., Zhang S., Lei S., Murage J., Fisk G.J.,
Li Y., Xu C., Pang R., Giegler K., Rao M.S., Mandalam R.,
Lebkowich J and Stanton L.W.
TITLE transcriptome characterization elucidates signaling networks that
control human ES cell growth and differentiation
JOURNAL Nat. Biotechnol. 22 (6), 707-716 (2004)
COMMENT Contact: Brandenberger R

Regenerative Medicine
Geron Corporation
230 Constitution Drive, Menlo Park, CA 94025, USA
Tel: 650 473 8658
Fax: 650 473 7760
Email: brandendberger@geron.com
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QY	106	AACCCGAGTTGCTTCGAGAGAGCGCTGTGTGAGCTTGAGGCTTCAACGATTACTTGTGAC	165	
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 Db 312 AACACCTTCTCCCTGGCTTGGAGTCTTACTGAGAGAGACTTACTACTACTGCTG 311
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 QY 466 AGGAGTCTGAGTCCAGGCTTCTGTTGGAGGCTCCAGAGAGTGGACATCCAGGCTG 525
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 QY 526 CGAGGGGGGAGACCTCCAGC 546
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 LOCUS CB96846
 DEFINITION AGENCOURT 1366148 NIH_MGC_148 Homo sapiens cDNA clone
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 VERSION CB96846
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 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 834)
 NIH-MGC http://mgs.nci.nih.gov/.
 National Institutes of Health.
 Unpublished (1999)
 Contact: Robert Strausberg, Ph.D.
 Email: cgaabs@mail.nih.gov
 Tissue Procurement: Dr. Stefan Hansson
 cDNA Library Preparation: Michael J. Brownstein (NHGRI) with help
 and advice from Piero Carninci (RIKEN)
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Found through the I.M.A.G.E. Consortium/LNL at:
 http://image.lnl.gov
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 and size 2.3 kb and normalized to 10^5. This is a primary
 library enriched for full-length clones and constructed
 using the Cap-trapper method (Carninci, in preparation).
 Library constructed by M. Brownstein (NHGRI, in preparation).
 National Institutes of Health). Note: this is a NIH_MGC
 Library."

ORIGIN

Query Match 91.2%; Score 498.2; DB 6; Length 834;
 Best Local Similarity 97.2%; Pred. No. 1,1e-117;
 Matches 528; Conservative 0; Mismatches 13; Indels 2; Gaps 2;

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 QY 181 TACGAGAGCGCGGCTCCCTGAGAGGCGCGGAGCGTTGCTTGTACATGTTGCTG 278
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 QY 361 GCTTGTGATCTTACTCTGAGAGAGACTTACTACTACTCTGCTGCTGCTGCTG 458
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 QY 480 AGCCCATCTCTGTTGGAGGCTT GAGAGAGTGGACATCCAGGCTGCGAGGGGGGAGA 578
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 Db 639 CAC 641

FEATURES

RESULT 10
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 VERSION CD671399.1 GI:2173139
 SOURCE EST.
 ORGANISM Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 592)
 Touchman, J.W., Bouffard, G., Smith, D., and Peterson, K.
 Project: steroid-response factors and similarities with retinal
 pigment epithelium
 Mol. Vis. 8 (4), 185-195 (2002)
 22103462
 12107412
 Contact: Mstow G
 Section on Molecular Structure and Function
 National Eye Institute
 6/331, NIH, Bethesda, MD 20892-2740, USA
 Tel: 301 402 3452

REFERENCE

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 TITLE Touchman, J.W., Bouffard, G., Smith, D., and Peterson, K.
 Project: steroid-response factors and similarities with retinal
 pigment epithelium
 JOURNAL Mol. Vis. 8 (4), 185-195 (2002)
 MEDLINE 22103462
 PUBMED 12107412
 COMMENT Contact: Mstow G
 Section on Molecular Structure and Function
 National Eye Institute
 6/331, NIH, Bethesda, MD 20892-2740, USA
 Tel: 301 402 3452

Fax: 301 496 0078
Email: greame@helix.nih.gov
Plate: 04 row: c column: 02
Seq primer: M13RPI reverse primer (ABI).
Location/Qualifiers
Source
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ORIGIN

Query Match 91.1%; Score 497.4; DB 6; Length 592;
Best Local Similarity 99.8%; Pred. No. 1.7e-117;
Matches 498; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

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(without alignments)
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Gapop 10.0 , Gapext 0.5

Searched: 1421835 seqs, 332370683 residues

Total number of hits satisfying chosen parameters: 3

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Listing first 1500 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	989	100.0	201	17	US-10-698-907-12
3	971	98.2	201	13	US-10-138-787-8

ALIGNMENTS

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; Sequence 4, Application US/09904954
; Patent No. US20020010325A1
; GENERAL INFORMATION:

APPLICANT: BECKMANN, M. P.
CERRETTI, DOUGLAS P.
TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE RECEPTOR HEK
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: IMMUNEX CORPORATION
STREET: 51 UNIVERSITY STREET
CITY: SEATTLE
STATE: WASHINGTON
COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/904,954
FILING DATE: 12-Jul-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/240,124
FILING DATE: <Unknown>
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESH, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 201 amino acids
TYPE: amino acid
MOLECULE TYPE: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 4:
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DB 61 YEGGPEGPEPTFLVYWDMPGYESCOAEGPRAYKRWCSLPFGHVFSEKIQFTFSL 120
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RESULT 2
US-10-698-907-12
; Sequence 12, Application US/10698907
; Publication No. US20050049194A1
; GENERAL INFORMATION:
; APPLICANT: Pilsen, Jonas
; APPLICANT: Holmberg, Johan
; TITLE OF INVENTION: Use of Ephrins and Related Molecules to Regulate Cellular
; FILE REFERENCE: 21882-529 UTIL
; CURRENT APPLICATION NUMBER: US/10/698,907

;; CURRENT FILING DATE: 2003-10-31
;; PRIOR APPLICATION NUMBER: US 60/460,488
;; PRIOR FILING DATE: 2003-04-03
;; PRIOR APPLICATION NUMBER: US 10/291,290
;; PRIOR FILING DATE: 2002-11-08
;; PRIOR APPLICATION NUMBER: US 60/393,272
;; PRIOR FILING DATE: 2002-07-02
;; PRIOR APPLICATION NUMBER: US 60/345,206
;; NUMBER OF SEQ ID NOS: 25
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;; SEQ ID NO 12
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;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-10-698-907-12

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RESULT 3

US-10-138-787-8
;; Sequence 8, Application US/10138787
;; Publication No. US20020172984A1
;; GENERAL INFORMATION:
;; APPLICANT: Holland, Sacha
;; APPLICANT: Mbamalu, Geraldine
;; APPLICANT: Pawson, Tony
;; TITLE OF INVENTION: OLIGOMERIZED RECEPTORS WHICH AFFECT PATHWAYS REGULATED
;; TITLE OF INVENTION: BY TRANSMEMBRANE LIGANDS FOR ELK-RELATED RECEPTOR
;; FILE REFERENCE: 11757.23USWO
;; CURRENT APPLICATION NUMBER: US/10/138,787
;; CURRENT FILING DATE: 2002-05-03
;; PRIOR APPLICATION NUMBER: US/09/214,631
;; PRIOR FILING DATE: 1999-03-12
;; PRIOR APPLICATION NUMBER: PCT/CA97/00473
;; PRIOR FILING DATE: 1997-07-04
;; PRIOR APPLICATION NUMBER: 60/021,272
;; NUMBER OF SEQ ID NOS: 13
;; SOFTWARE: PatentIn Ver. 2.0
;; SEQ ID NO 8
;; LENGTH: 201
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-10-138-787-8

Query Match
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Matches 176; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

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Maximum DB seq length: 200000000

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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2	989	100.0	201	1	US-08-453-943-4 Sequence 4, Appli
3	989	100.0	201	2	US-09-057-121-4 Sequence 4, Appli
4	989	100.0	201	3	US-09-358-734-4 Sequence 4, Appli
5	989	100.0	210	4	US-09-949-016-7162 Sequence 7162, Ap
6	971	98.2	201	4	US-09-214-631-8 Sequence 8, Appli

ALIGNMENTS

RESULT 1
US-08-240-124-4
; Sequence 4, Application US/08240124
; Patent No. 5516658
; GENERAL INFORMATION:
; APPLICANT: BECKMANN, M. P.
; APPLICANT: CERRETTI, DOUGLAS P.
; TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
; TITLE OF INVENTION: RECEPTOR HEK
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: IMMUNEX CORPORATION
; STREET: 51 UNIVERSITY STREET
; CITY: SEATTLE
; STATE: WASHINGTON
; COUNTRY: USA

ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/240,124
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/161,132
FILING DATE: 03-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
TELEX: 756822
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 201 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-240-124-4

Query Match 100.0%; Score 989; DB 1; Length 201;
Best Local Similarity 100.0%; Pred. No. 2.6e-107;
Matches 179; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	MRLPLRLRTVLMMAFLGSP	SRHVVYNNSSNRLRGDAVVELGNDYLDIVCPH	60
DB	1	MRLPLRLRTVLMMAFLGSP	SRHVVYNNSSNRLRGDAVVELGNDYLDIVCPH	60
QY	61	YEGPGPEGETALVWMPGVESCOAEGPRAYKXWVCSLPGHQVPSKIRFTPESL		120
DB	61	YEGPGPEGETALVWMPGVESCOAEGPRAYKXWVCSLPGHQVPSKIRFTPESL		120
QY	121	GFEFLPGEITYYISVTPRESSGQCLRLQVSVCKEKRESAHPVGSRGSGTSGMRGCD		179
DB	121	GFEFLPGEITYYISVTPRESSGQCLRLQVSVCKEKRESAHPVGSRGSGTSGMRGCD		179

RESULT 2
US-08-453-943-4
; Sequence 4, Application US/08453943
; Patent No. 5738844
; GENERAL INFORMATION:
; APPLICANT: BECKMANN, M. P.
; APPLICANT: CERRETTI, DOUGLAS P.
; TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
; TITLE OF INVENTION: RECEPTOR HEK
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: IMMUNEX CORPORATION
; STREET: 51 UNIVERSITY STREET
; CITY: SEATTLE
; STATE: WASHINGTON
; COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Apple System 7.1

SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/453,943
FILING DATE: 30-MAY-1995
CLASSIFICATION: 510
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/240,124
FILING DATE: 09-MAY-1994
APPLICATION NUMBER: US 08/161,132
FILING DATE: 03-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 201 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-453-943-4

Query Match
Best Local Similarity 100.0%; Score 989; DB 1; Length 201;
Matches 179; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRLPLRLTYMAAFGLSPRLRGSSLRHYVYWNSSNPRLRGDAVVELGLNDYLDIVCPH 60
DB 1 MRLPLRLTYMAAFGLSPRLRGSSLRHYVYWNSSNPRLRGDAVVELGLNDYLDIVCPH 60
QY 61 YEGGPPPEGPEFTFALYVMDMPGYSCQAEGRPRAYRWVCSLPFGHVQFSEKIQRTPTPSL 120
DB 61 YEGGPPPEGPEFTFALYVMDMPGYSCQAEGRPRAYRWVCSLPFGHVQFSEKIQRTPTPSL 120
QY 121 GFEFLPGETYYIISVFTPESSGQCLRLQVSVCKEKRSSEAHFVSGSPGSGTSGMRGCD 179
DB 121 GFEFLPGETYYIISVFTPESSGQCLRLQVSVCKEKRSSEAHFVSGSPGSGTSGMRGCD 179

RESULT 3

US-09-057-121-4
Sequence 4, Application US/09057121
Patent No. 5969110
GENERAL INFORMATION:
APPLICANT: BECKMANN, M. P.
TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: IMMUNEX CORPORATION
STREET: 51 UNIVERSITY STREET
CITY: SEATTLE
STATE: WASHINGTON
COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: Apple Macintosh
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/057,121

FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/240,124
FILING DATE:
APPLICATION NUMBER: US 08/161,132
FILING DATE: 03-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 201 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-057-121-4

Query Match
Best Local Similarity 100.0%; Score 989; DB 2; Length 201;
Matches 179; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRLPLRLTYMAAFGLSPRLRGSSLRHYVYWNSSNPRLRGDAVVELGLNDYLDIVCPH 60
DB 1 MRLPLRLTYMAAFGLSPRLRGSSLRHYVYWNSSNPRLRGDAVVELGLNDYLDIVCPH 60
QY 61 YEGGPPPEGPEFTFALYVMDMPGYSCQAEGRPRAYRWVCSLPFGHVQFSEKIQRTPTPSL 120
DB 61 YEGGPPPEGPEFTFALYVMDMPGYSCQAEGRPRAYRWVCSLPFGHVQFSEKIQRTPTPSL 120
QY 121 GFEFLPGETYYIISVFTPESSGQCLRLQVSVCKEKRSSEAHFVSGSPGSGTSGMRGCD 179
DB 121 GFEFLPGETYYIISVFTPESSGQCLRLQVSVCKEKRSSEAHFVSGSPGSGTSGMRGCD 179

RESULT 4

US-09-358-734-4
Sequence 4, Application US/09358734
Patent No. 6274117
GENERAL INFORMATION:
APPLICANT: BECKMANN, M. P.
TITLE OF INVENTION: CYTOKINE THAT BINDS THE CELL SURFACE
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: IMMUNEX CORPORATION
STREET: 51 UNIVERSITY STREET
CITY: SEATTLE
STATE: WASHINGTON
COUNTRY: USA
ZIP: 98101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: Apple Macintosh
SOFTWARE: Microsoft Word for Apple, Version 5.1a
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/358,734
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/240,124
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/114,426
FILING DATE: 30-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/109,745
FILING DATE: 20-AUG-1993
ATTORNEY/AGENT INFORMATION:
NAME: SEESE, KATHRYN A.
REGISTRATION NUMBER: 32,172
REFERENCE/DOCKET NUMBER: 2814-C
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 587-0430
TELEFAX: (206) 233-0644
TELEX: 756822
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 201 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-358-734-4

Query Match 100.0%; Score 989; DB 3; Length 201;
Best Local Similarity 100.0%; Pred. No. 2.6e-107;
Matches 179; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 60
DB 1 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 60
QY 61 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 120
DB 61 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 120
QY 121 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 179
DB 121 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 179

RESULT 5
US-09-949-016-7162
Sequence 7162, Application US/09949016
Patent No. 6812339
GENERAL INFORMATION:
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 7162
LENGTH: 210
TYPE: PRT
ORGANISM: Human
US-09-949-016-7162

Query Match 100.0%; Score 989; DB 4; Length 210;
Best Local Similarity 100.0%; Pred. No. 2.8e-107;
Matches 179; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 60
DB 10 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 69

QY 61 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 120
DB 70 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 129
QY 121 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 179
DB 130 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 188

RESULT 6
US-09-214-631-8
Sequence 8, Application US/09214631
Patent No. 6413730
GENERAL INFORMATION:
APPLICANT: Holland, Sacha
APPLICANT: Mamanu, Geraldine
APPLICANT: Pawson, Tony
TITLE OF INVENTION: OLIGOMERIZED RECEPTORS WHICH AFFECT PATHWAYS REGULATED
BY TRANSMEMBRANE LIGANDS FOR ELK-RELATED RECEPTOR
TITLE OF INVENTION: TYROSINE KINASES
FILE REFERENCE: 11757.23USWO
CURRENT APPLICATION NUMBER: US/09/214,631
CURRENT FILING DATE: 1999-03-12
EARLIER APPLICATION NUMBER: PCT/CA97/00473
EARLIER FILING DATE: 1997-07-04
EARLIER APPLICATION NUMBER: 60/021,272
EARLIER FILING DATE: 1996-07-05
NUMBER OF SEQ ID NOS: 13
SOFTWARE: Patentln Ver. 2.0
SEQ ID NO 8
LENGTH: 201
TYPE: PRT
ORGANISM: Homo sapiens
US-09-214-631-8

Query Match 98.2%; Score 971; DB 4; Length 201;
Best Local Similarity 98.3%; Pred. No. 3.3e-105;
Matches 176; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 60
DB 1 MRLPLRLTYLMAAFGLGSPRLRGSSSLRHVVYWNSSNPRLLRGDAVVELGLNDYLDIVCPH 60
QY 61 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 120
DB 61 YEGGPPGPEPTFALYVNDMPGYESCOAEGPRAYKRWVCSLPFGHVQFSEKIQFTFPSL 120
QY 121 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 179
DB 121 GFELPGETYYIISVPTPSSGQCLRLQVSVCKERSBSAHFVGSFGESGTSGMRGSD 179

Search completed: April 19, 2005, 23:19:49
Job time: 14.1316 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using bw model

Run on: April 19, 2005, 09:30:04 ; Search time 344.062 Seconds
(without alignments)
8258.620 Million cell updates/sec

Title: US-09-904-954-3_COPY_94_573
Perfect score: 480
Sequence: 1 GGCTCCAGCCTCCGCGACACTCCAGC 480

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 7

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 89%
Maximum Match 100%
Listing first 1500 summaries

Database : N_Geneseq_16Dec04:*
1: geneseqn1980s:*
2: geneseqn1980s:*
3: geneseqn2000s:*
4: geneseqn2001s:*
5: geneseqn2001bs:*
6: geneseqn2002as:*
7: geneseqn2002bs:*
8: geneseqn2003as:*
9: geneseqn2003bs:*
10: geneseqn2003cs:*
11: geneseqn2003ds:*
12: geneseqn2004as:*
13: geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	480	100.0	636	2	AAQ85888 Human hek
2	480	100.0	1181	8	ABZ34863 Coding se
3	478.4	99.7	606	6	ABV78135 Human eph
4	478.4	99.7	606	6	ABZ35711 Human eph
5	478.4	99.7	606	6	ABX09954 Human eph
6	478.4	99.7	606	6	ABL91676 Human pol
7	432.8	90.2	770	12	ADN02769 Human rec

ALIGNMENTS

RESULT 1
AAQ85888
ID AAQ85888 standard; cDNA to mRNA; 636 BP.
XX
AC AAQ85888;
XX
DT 25-MAR-2003 (revised)
DT 03-OCT-1995 (first entry)

XX	Human hek-L protein cDNA clone C6.
DE	
XX	Ligand; cell surface; tyrosine kinase receptor; tumorigenesis; immunogen;
KW	88.
XX	
OS	Homo sapiens.
XX	
FT	Key CDS
FT	Location/Qualifiers
FT	28..633
FT	/*tag= a
FT	28..93
FT	/*tag= b
FT	94..630
FT	mat_peptide
FT	/*tag= c
XX	
XX	W09506065-A1.
XX	
XX	02-MAR-1995.
PD	
XX	
XX	17-AUG-1994; 94WO-US009282.
XX	
XX	20-AUG-1993; 93US-00109745.
PR	30-AUG-1993; 93US-00114426.
PR	03-DEC-1993; 93US-00161132.
PR	09-MAY-1994; 94US-00240124.
XX	
PA	(IMMV) IMMUNEX CORP.
XX	
XX	Beckmann MP, Cerretti DP;
PI	
XX	WPI; 1995-106811/14.
DR	P-PSDB; AAR71482.
DR	
XX	
PT	New isolated DNA encoding hek-L protein or its fusion products - useful
PT	as assay reagent or for carrying therapeutic and diagnostic compounds to
PT	leukaemia cells.
XX	
PS	Claim 3; Page 37; 45pp; English.
XX	
XX	The sequence is that of a clone encoding hek-L protein, a protein that
CC	can bind hek (a cell surface receptor tyrosine kinase). Hek-L is the
CC	first known ligand for hek and can be used to study cellular processes
CC	regulated by hek (which may be involved in tumorigenesis). It is also an
CC	immunogen for antibody production, as a reagent for detecting hek or hek-
CC	L in vitro assays, to determine binding of hek proteins, to purify hek
CC	proteins, and to carry diagnostic or cytotoxic agents to particular
CC	leukaemia cells that express the hek antigen. Hek-L also binds the elk
CC	tyrosine kinase receptors. See also AAQ85887. (Updated on 25-MAR-2003 to
CC	correct PN field.)
XX	
SQ	Sequence 636 BP; 102 A; 202 C; 186 G; 146 T; 0 U; 0 Other;
XX	
XX	
XX	Query Match 100.0%; Score 480; DB 2; Length 636;
XX	Best Local Similarity 100.0%; Pred. No. 2,2e-129; Indels 0; Gaps 0;
XX	Matches 480; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 GGCTCCAGCCTCCGCGACACTCCAGTGAACCTCCAGTGAACCCAGGTTGCTTGAAGA 60
DB	94 GGCTCCAGCCTCCGCGACACTCCAGTGAACCTCCAGTGAACCCAGGTTGCTTGAAGA 153
QY	61 GAGCCCGTGTGAGAGCTGGGCTTCAGATTTACTGTAACATTTGTCTGCCCTCAAGAA 120
DB	154 GAGCCCGTGTGAGAGCTGGGCTTCAGATTTACTGTAACATTTGTCTGCCCTCAAGAA 213
QY	121 GGCCAGAGGCCCCCTGAGGCCCCGAGACCTTTGCTTTGTACATGTGAGACTGGCCAGGC 180
DB	214 GGCCAGAGGCCCCCTGAGGCCCCGAGACCTTTGCTTTGTACATGTGAGACTGGCCAGGC 273
QY	181 TATGAGTCTTGCCAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 240
DB	274 TATGAGTCTTGCCAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 333

CC both in vivo and in vitro and also increases stability and thus the
CC effective concentration inside the cell. The present sequence is that of
CC a gene related to the invention
XX

Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match 99.7%; Score 478.4; DB 6; Length 606;
Best Local Similarity 99.8%; Pred. No. 6,4e-129;
Matches 479; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

```
QY 1 GGGTCAGAGCTCCGCGCAGTGTCTACTGAGAACTCCGAGTCCAGAGTCTTCCAGGA 60
DB 67 GGGTCAGAGCTCCGCGCAGTGTCTACTGAGAACTCCGAGTCCAGAGTCTTCCAGGA 126
QY 61 GAGCGGTGTGTGAGAGTGTGAGTCTCAAGATTACTAGAACTGTCTGCCCCCACTAGAA 120
DB 127 GAGCGGTGTGTGAGAGTGTGAGTCTCAAGATTACTAGAACTGTCTGCCCCCACTAGAA 186
QY 121 GGGCCAGAGGCCCCCTGAGAGGCCCCGAGACGTTTGTCTTGTACATGTGACTGAGCCAGGC 180
DB 187 GGGCCAGAGGCCCCCTGAGAGGCCCCGAGACGTTTGTCTTGTACATGTGACTGAGCCAGGC 246
QY 181 TATGAGTCTGTCAGAGGAGGAGGCCCCGAGGCTTACAGGCGTGGGTGTGCTCCGAGCC 240
DB 247 TATGAGTCTGTCAGAGGAGGAGGCCCCGAGGCTTACAGGCGTGGGTGTGCTCCGAGCC 306
QY 241 TTTGGCCATGTTCAATTCTCAGAGAAATTCAGAGGCTTCAACCTTCTCCCTCGGCTTT 300
DB 307 TTTGGCCATGTTCAATTCTCAGAGAAATTCAGAGGCTTCAACCTTCTCCCTCGGCTTT 366
QY 301 GAGTCTTACCTGAGAGACTTACTACTACATCTCGGTGCCACTCCAGAGATTCTGGC 360
DB 367 GAGTCTTACCTGAGAGACTTACTACTACATCTCGGTGCCACTCCAGAGATTCTGGC 426
QY 361 CAGTCTTGAAGGCTCCAGGTGTCTGTCTGTGCAAGAGAGAAAGTGTGAGTCAAGCCAT 420
DB 427 CAGTCTTGAAGGCTCCAGGTGTCTGTCTGTGCAAGAGAGAAAGTGTGAGTCAAGCCAT 486
QY 421 CCGTGTGGAGGCCCCCTGAGAGAGTGGCAATCAGAGGTGGAGAGGGGGGACACTCCAGC 480
DB 487 CCGTGTGGAGGCCCCCTGAGAGAGTGGCAATCAGAGGTGGAGAGGGGGGACACTCCAGC 546
```

RESULT 4

ABZ35711 standard; DNA; 606 BP.

ABZ35711;

07-FEB-2003 (first entry)

Human ephrin A3 encoding polynucleotide SEQ ID NO 19.

Double stranded RNA; dsRNA; RNAi; RNA inhibition; cytosolic; virucide;
KM protozoicide; gene expression; antisense; tumor; infection; Plasmodium;
KM virus; viroid; anti-GFP; human; HIV; human immunodeficiency virus;
KM Hepatitis B virus; human papilloma virus; gene; ds.

Homo sapiens.

DE10100588-A1.

18-JUL-2002.

09-JAN-2001; 2001DE-01000588.

09-JAN-2001; 2001DE-01000588.

(RIBO-) RIBOPHARMA AG.

Kreutzer R, Limmer S, Rost S, Hadwiger P;

WPI; 2002-683450/74.

XX Inhibiting expression of target genes, useful e.g. for treating tumors,
PT by introducing into cells two double-stranded RNAs that are complementary
PT to the target.

PS Claim 13; Page 19-20; 100pp; German.

XX The invention relates to inhibiting expression of a target gene in a cell
CC by introducing at least two oligonucleotides (dsRNA and i), both
CC with a double-stranded (ds) structure of at most 49 sequential nucleotide
CC pairs. At least part of one strand (S1, S2) of the ds structures in each
CC of dsRNA and i are complementary to regions in the target gene. The
CC method uses antisense inhibition of gene expression using double stranded
CC RNA inhibition (RNAi). The method is particularly used to treat tumors
CC or infections, especially by Plasmodium or viruses/viroids (pathogenic on
CC humans, animals or plants). The method provides more effective inhibition
CC of expression than known methods using a single dsRNA, even at very low
CC concentrations. When dsRNA has at least one unpaired nucleotide at the
CC end, stability (and thus effective concentration in the cell) is improved
CC and efficiency can be increased further by pretreating the cells with
CC interferon. The present sequence is that of a target DNA of the invention

Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match 99.7%; Score 478.4; DB 6; Length 606;
Best Local Similarity 99.8%; Pred. No. 6,4e-129;
Matches 479; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 1 GGGTCAGAGCTCCGCGCAGTGTCTACTGAGAACTCCGAGTCCAGAGTCTTCCAGGA 60
DB 67 GGGTCAGAGCTCCGCGCAGTGTCTACTGAGAACTCCGAGTCCAGAGTCTTCCAGGA 126
QY 61 GAGCGGTGTGTGAGAGTGTGAGTCTCAAGATTACTAGAACTGTCTGCCCCCACTAGAA 120
DB 127 GAGCGGTGTGTGAGAGTGTGAGTCTCAAGATTACTAGAACTGTCTGCCCCCACTAGAA 186
QY 121 GGGCCAGAGGCCCCCTGAGAGGCCCCGAGACGTTTGTCTTGTACATGTGACTGAGCCAGGC 180
DB 187 GGGCCAGAGGCCCCCTGAGAGGCCCCGAGACGTTTGTCTTGTACATGTGACTGAGCCAGGC 246
QY 181 TATGAGTCTGTCAGAGGAGGAGGCCCCGAGGCTTACAGGCGTGGGTGTGCTCCGAGCC 240
DB 247 TATGAGTCTGTCAGAGGAGGAGGCCCCGAGGCTTACAGGCGTGGGTGTGCTCCGAGCC 306
QY 241 TTTGGCCATGTTCAATTCTCAGAGAAATTCAGAGGCTTCAACCTTCTCCCTCGGCTTT 300
DB 307 TTTGGCCATGTTCAATTCTCAGAGAAATTCAGAGGCTTCAACCTTCTCCCTCGGCTTT 366
QY 301 GAGTCTTACCTGAGAGACTTACTACTACATCTCGGTGCCACTCCAGAGATTCTGGC 360
DB 367 GAGTCTTACCTGAGAGACTTACTACTACATCTCGGTGCCACTCCAGAGATTCTGGC 426
QY 361 CAGTCTTGAAGGCTCCAGGTGTCTGTCTGTGCAAGAGAGAAAGTGTGAGTCAAGCCAT 420
DB 427 CAGTCTTGAAGGCTCCAGGTGTCTGTCTGTGCAAGAGAGAAAGTGTGAGTCAAGCCAT 486
QY 421 CCGTGTGGAGGCCCCCTGAGAGAGTGGCAATCAGAGGTGGAGAGGGGGGACACTCCAGC 480
DB 487 CCGTGTGGAGGCCCCCTGAGAGAGTGGCAATCAGAGGTGGAGAGGGGGGACACTCCAGC 546
```

RESULT 5

ABX09954 standard; DNA; 606 BP.

ABX09954;

23-JAN-2003 (first entry)

Human ephrin A3 DNA fragment SEQ ID 19.

Oligonucleotide; interferon; oncogene; cytokine; Id; developmental;
KM prion; inhibition; human; ds.

XX OS Homo sapiens.
 XX PN DE10100587-C1.
 XX PD 21-NOV-2002.
 XX PF 09-JAN-2001; 2001DE-01000587.
 XX PR 09-JAN-2001; 2001DE-01000587.
 XX PA (RIBO-) RIBOPHARMA AG.
 XX PI Kreutzer R, Limmer S, Rost S, Hadwiger P;
 XX DR MPI, 2002-742209/81.
 XX PT Inhibiting expression of target genes, e.g. oncogenes, in cells, by
 XX PT introducing of complementary double-stranded oligoribonucleotide, after
 XX PT treating the cell with interferon.
 XX PS Disclosure; Page 24-25; 98pp; German.

CC This invention describes a novel method for inhibiting expression of a
 CC target gene by introducing into the cell that contains the target gene at
 CC least one oligoribonucleotide (dsRNA) that has a double-stranded (ds)
 CC structure of not more than 49 consecutive nucleotides (nt), where at
 CC least a segment of one strand of the ds structure is complementary with
 CC the target gene and the cells are treated with interferon before
 CC introduction of dsRNA. The method is used to inhibit expression of
 CC target genes, particularly oncogenes, cytokine genes, Id (not defined)
 CC protein genes, developmental or prion genes, or genes expressed in
 CC pathogenic organisms (particularly plasmids) or in viruses or viroids
 CC (pathogenic in humans, animals or plants). Treating the cells with
 CC interferon greatly increases the extent to which dsRNA can inhibit
 CC expression of the target genes, and the effect is even greater when dsRNA
 CC are modified to increase their stability. ABX09336-ABX10075 represent
 CC gene fragments used to illustrate the method of the invention

Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

Query Match

Best Local Similarity 99.7%; Score 478.4; DB 6; Length 606;
 Matches 479; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGCTCCAGGCTCCGCGCAGTCTGAACTGCAATCCCAAGGTTGCTTCAGGA 60
 DB 67 GGCTCCAGGCTCCGCGCAGTCTGAACTGCAATCCCAAGGTTGCTTCAGGA 60
 QY 61 GAGCCGTGTGAGAGCTGGGCTCAACGATTACCTAGACATTGCTGCCCCCACTAGAA 126
 DB 127 GAGCCGTGTGAGAGCTGGGCTCAACGATTACCTAGACATTGCTGCCCCCACTAGAA 126
 QY 121 GGCCCAAGGCCCCCGAGGCGCCGAGACGTTGCTTGTACATGAGTGAAGTGGCCAGGC 180
 DB 187 GGCCCAAGGCCCCCGAGGCGCCGAGACGTTGCTTGTACATGAGTGAAGTGGCCAGGC 180
 QY 181 TATGAGTCTGCGCAGGAGAGGCGCCCGGAGCTCAAGAGGCTGGGTGCTTCCTGCCC 240
 DB 247 TATGAGTCTGCGCAGGAGAGGCGCCCGGAGCTCAAGAGGCTGGGTGCTTCCTGCCC 240
 QY 241 TTGGCCATGTTCAATTCTCAGAGAAATTCAAGCGGTTCAACCTTTCCCTCGGCTTT 300
 DB 307 TTGGCCATGTTCAATTCTCAGAGAAATTCAAGCGGTTCAACCTTTCCCTCGGCTTT 300
 QY 301 GAGTTTCTTACCTGAGAGACTTACTACTCATCTGGTGGCCCACTCCAGAGATTGCGC 360
 DB 367 GAGTTTCTTACCTGAGAGACTTACTACTCATCTGGTGGCCCACTCCAGAGATTGCGC 360
 QY 361 CAGTGTGAGAGCTCCAGGAGTGTCTGCTCTGCAAGAGAGAAATGAGTCAAGCCAT 420
 DB 427 CAGTGTGAGAGCTCCAGGAGTGTCTGCTCTGCAAGAGAGAAATGAGTCAAGCCAT 420

QY 421 CCTGTGGAGGCTTCGAGAGAGTGGCAGATCAGAGTGGCGAGGCGGAGCACTCCAGC 480
 DB 487 CCTGTGGAGGCTTCGAGAGAGTGGCAGATCAGAGTGGCGAGGCGGAGCACTCCAGC 546

RESULT 6

ID ABL91676 standard; DNA; 606 BP.

XX ABL91676;

XX 28-MAY-2002 (first entry)

DE Human polynucleotide SEQ ID NO 19.

XX Human; HIV; HCV; gene expression; oligoribonucleotide; tumour; pathogen;

XX Plasmodium; virus; viroid; cytokine; prion; antisense oligonucleotide;

XX cytosolic; virucide; protozoacide; antibacterial; ds.

OS Homo sapiens.

XX DE10100586-C1.

XX 11-APR-2002.

XX 09-JAN-2001; 2001DE-01000586.

XX 09-JAN-2001; 2001DE-01000586.

XX (RIBO-) RIBOPHARMA AG.

XX Kreutzer R, Limmer S, Rost S, Hadwiger P;

XX MPI, 2002-270454/32.

XX Claim 13; Page 20-21; 104pp; German.

CC The invention relates to a method for inhibiting expression of a target

CC gene (ABL91658-ABL91797) in a cell by introducing at least one

CC oligoribonucleotide that has a double-stranded structure consisting of at

CC most 49 sequential nucleotide pairs, with at least part of one strand

CC complementary with the target gene and has at least one end a single-

CC stranded segment of 1-4 nt. The method provides oligoribonucleotides for

CC but the oligoribonucleotides may also be directed against genes present

CC in pathogens (e.g. Plasmodium or viruses/viroids), pathogenic on humans,

CC animals or plants) or against cytokine, Id, developmental or prion genes.

CC The method provides more effective inhibition of gene expression than use

CC of known oligonucleotides, probably because the unpaired overhang

CC increases stability and thus intracellular concentration

XX Sequence 606 BP; 96 A; 191 C; 175 G; 144 T; 0 U; 0 Other;

QY Query Match 99.7%; Score 478.4; DB 6; Length 606;
 Best Local Similarity 99.8%; Pred. No. 6.4e-129;
 Matches 479; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GGCTCCAGGCTCCGCGCAGTCTGAACTGCAATCCCAAGGTTGCTTCAGGA 60
 DB 67 GGCTCCAGGCTCCGCGCAGTCTGAACTGCAATCCCAAGGTTGCTTCAGGA 60
 QY 61 GAGCCGTGTGAGAGCTGGGCTCAACGATTACCTAGACATTGCTGCCCCCACTAGAA 120
 DB 127 GAGCCGTGTGAGAGCTGGGCTCAACGATTACCTAGACATTGCTGCCCCCACTAGAA 120
 QY 121 GGCCCAAGGCCCCCGAGGCGCCGAGACGTTGCTTGTACATGAGTGAAGTGGCCAGGC 180
 DB 187 GGCCCAAGGCCCCCGAGGCGCCGAGACGTTGCTTGTACATGAGTGAAGTGGCCAGGC 180

QY 181 TATGAGTCTGCGAGAGAGGCCCCGGGCTTACAGGCGTGGTGTGCTCCCTGCC 240
 DB 247 TATGAGTCTGCGAGAGAGGCCCCGGGCTTACAGGCGTGGTGTGCTCCCTGCC 306
 QY 241 TTTGGCCATGTTCAATTCAGAGAGATTCAAGGCTTCAACCTTTCTCCCTGGCTTT 300
 DB 307 TTTGGCCATGTTCAATTCAGAGAGATTCAAGGCTTCAACCTTTCTCCCTGGCTTT 366
 QY 301 GAGTTCTTACCTGAGAGACTTACTACTACTCTCGGTGCCCATCCAGAGAGTTTGGC 360
 DB 367 GAGTTCTTACCTGAGAGACTTACTACTACTCTCGGTGCCCATCCAGAGAGTTTGGC 426
 QY 361 CAGTCTTGAAGGCTCCAGGTCTGTCTGTCTGCAAGAGAGAGAGTCTGAGTCAAGCCAT 420
 DB 427 CAGTCTTGAAGGCTCCAGGTCTGTCTGTCTGCAAGAGAGAGAGTCTGAGTCAAGCCAT 486
 QY 421 CCGTGTGGAGAGCCCTGAGAGAGTGGACATCAGGGTGGCGAGGGGGAGACATCCGAGC 480
 DB 487 CCGTGTGGAGAGCCCTGAGAGAGTGGACATCAGGGTGGCGAGGGGGAGACATCCGAGC 546

RESULT 7
 ADN02769
 ID ADN02769 standard; cDNA; 770 BP.
 XX
 AC ADN02769;
 DT 01-JUL-2004 (first entry)
 XX
 DB Human receptor and membrane-associated protein cDNA #22.
 XX
 KW db; gene: cytosolic; antiarteriosclerotic; anti-HIV; cerebroprotective;
 KW antiParkinsonian; nootropic; neuroprotective; immunosuppressive;
 KW antiinflammatory; antiallergic; anabolic; hypertensive; anorectic;
 KW antidiabetic; gene therapy; receptor and membrane-associated protein;
 KW REMAP; diagnosing; cancer; atherosclerosis; AIDS; allergy;
 KW Parkinson's disease; Alzheimer's disease; stroke; Addison's disease;
 KW obesity; diabetes; microarray; gene expression; receptor.
 XX
 OS Homo sapiens.
 XX
 PN WO2004029218-A2.
 XX
 PD 08-APR-2004.
 XX
 PF 26-SEP-2003; 2003WO-US030894.
 XX
 PR 27-SEP-2002; 2002US-0414302P.
 PR 11-OCT-2002; 2002US-0417797P.
 PR 16-OCT-2002; 2002US-0419217P.
 PR 29-OCT-2002; 2002US-0422375P.
 PR 07-NOV-2002; 2002US-0424799P.
 XX
 PA (INCY-) INCYTE CORP.
 XX
 PI Marguis JP, Tran UK, Lee SY, Richardson TW, Chawla NK,
 PI Hafalia AJA, Becha SD, Ramkumar J, Khare R, Tang YT, Yue H,
 PI Baughn ME, Elliott VS, Swarnakar A, Lu DM, Policky JL,
 PI Thangavelu K, Gietzen KJ, Blake UJ, Ison CH,
 DR WPI; 2004-316100/29.
 DR P-PSDB; ADN027719.
 XX
 PT New human receptors and membrane-associated proteins and polynucleotides
 PT for diagnosing, preventing or treating diseases associated with aberrant
 PT protein expression, e.g. cancer, atherosclerosis, AIDS, stroke or
 PT diabetes.
 XX
 PS Claim 5; SEQ ID NO 72; 268bp; English.
 XX
 CC The invention relates to novel human receptor and membrane-associated
 CC protein (REMAP) and the genes encoding them, a naturally-occurring amino
 CC acid sequence that is at least 90% to at least 98% identical any of the

CC amino acid sequences cited above; or a biologically active or immunogenic
 CC fragment of the polypeptide. The specification also discloses a naturally
 CC occurring polynucleotide sequence that is at least 90% to at least 98%
 CC identical to any of the nucleotide sequences cited above; their
 CC complements or an RNA equivalent. The composition and methods are useful
 CC for diagnosing, preventing or treating diseases or conditions associated
 CC with aberrant expression of REMAP, such as cell proliferative (e.g.
 CC cancer or atherosclerosis), autoimmune/inflammatory (e.g. AIDS or
 CC allergies), neurological (e.g. Parkinson's disease, Alzheimer's disease
 CC or stroke), metabolic (e.g. Addison's disease or obesity), developmental
 CC or endocrine (e.g. diabetes) disorders. These may also be used for
 CC assessing the effects of exogenous compounds on the expression of nucleic
 CC acid and amino acid sequences of REMAP. The REMAP or its fragments are
 CC also useful in screening compounds for effectiveness as agonist or
 CC antagonist of the polypeptides, or in altering the expression of the
 CC target polynucleotide and compounds that specifically bind to or modulate
 CC the activity of the polypeptide. The microarray is useful in monitoring
 CC or measuring protein-protein interactions, drug-target interactions, and
 CC gene expression profiles. This sequence corresponds to the cDNA encoding
 CC one of the proteins of the invention.

XX Sequence 770 BP; 134 A; 242 C; 213 G; 181 T; 0 U; 0 Other;
 SQ
 Query Match 90.2%; Score 432.8; DB 12; Length 770;
 Best Local Similarity 98.4%; Pred. No. 1.3e-115;
 Matches 437; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 37 AGTAACCCAGGTTGCTTTCAGAGAGAGCCGCTGTGAGTGGGCTCAACGATTACTTA 96
 DB 252 AGTCTGCGAGGTTGCTTTCAGAGAGAGCCGCTGTGAGTGGGCTCAACGATTACTTA 311
 QY 97 GACATTGTCTGCCCCCACTACGAAAGGCCAGGGCCCCCTAGAGGCCCGGAGCTTTGCT 156
 DB 312 GACATTGTCTGCCCCCACTACGAAAGGCCAGGGCCCCCTAGAGGCCCGGAGCTTTGCT 371
 QY 157 TTGACATGCTGAGCTGGCCAGGCTATGAGTCTGTCGACGAGAGAGGCCCGGAGCTTAC 216
 DB 372 TTGACATGCTGAGCTGGCCAGGCTATGAGTCTGTCGACGAGAGAGGCCCGGAGCTTAC 431
 QY 217 AAGGCTGGGTGTGCTCCCTGCTTGGCCATGTTCAATTCAGAGAGATTAGCGC 276
 DB 432 AAGGCTGGGTGTGCTCCCTGCTTGGCCATGTTCAATTCAGAGAGATTAGCGC 491
 QY 277 TTCAACCTTTTCCTCGGCTTTGAGTTCTTACCTGGAAGACTTACTACTCTCG 336
 DB 492 TTCAACCTTTTCCTCGGCTTTGAGTTCTTACCTGGAAGACTTACTACTCTCG 551
 QY 337 GTGCCACTCCAGAGAGTTCTGGCCAGTGTCTTGGAGGCTCCAGAGTGTCTGCTGCAAG 396
 DB 552 GTGCCACTCCAGAGAGTTCTGGCCAGTGTCTTGGAGGCTCCAGAGTGTCTGCTGCAAG 611
 QY 397 GAGAGGAAGTCTGAGTCAAGCCATCTGTTGGAGGCCCTGAGAGAGTGGCACTACAGG 456
 DB 612 GAGAGGAAGTCTGAGTCAAGCCATCTGTTGGAGGCCCTGAGAGAGTGGCACTACAGG 671
 QY 457 TGGCGAGGGGGGAGCACTCCAGC 480
 DB 672 TGGCGAGGGGGGAGCACTCCAGC 695

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